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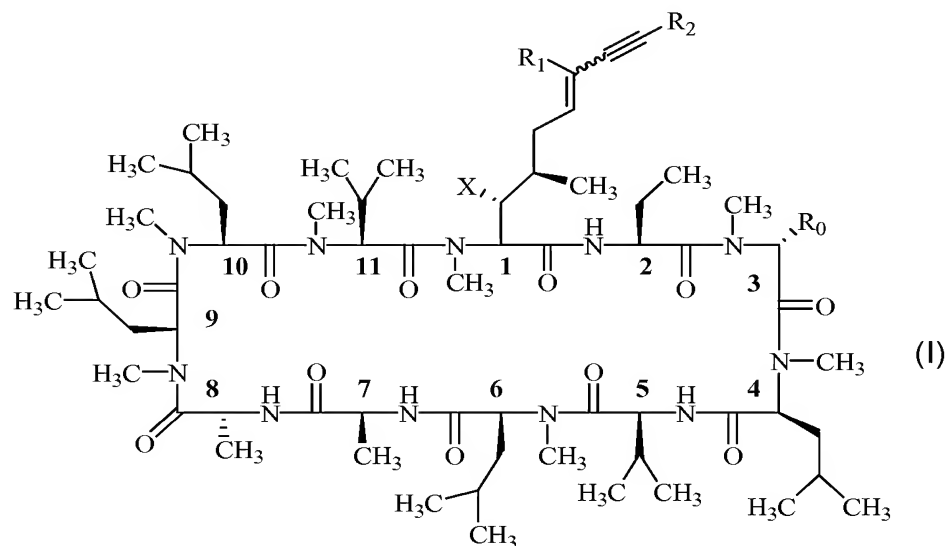
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(54) Title: USE OF CYCLOSPORIN ALKYNE/ALKENE ANALOGUES FOR PREVENTING OR TREATING VIRAL-INDUCED DISORDERS



(57) Abstract: The present invention relates to methods of preventing or treating a mammal with a viral-induced disorder. The method involves administering to the mammal a therapeutically effective amount of a compound represented by Formula I, as shown below: or a pharmaceutically acceptable salt thereof, with X, Ro, R1, and R2 defined herein, under conditions effective to prevent or treat the viral-induced disorder.

USE OF CYCLOSPORIN ALKYNE/ALKENE ANALOGUES FOR PREVENTING OR TREATING VIRAL-INDUCED DISORDERS

[0001] This application claims the benefit of U.S. Patent Application Serial
5 No. 11/391,023, filed March 28, 2006, which is hereby incorporated by reference in its entirety.

FIELD OF THE INVENTION

[0002] The present invention discloses novel cyclosporin alkyne/alkene
10 analogues and their utilities as pharmaceutical agents for prevention and treatment of viral-induced diseases. Methods for preparation of such analogues are also disclosed.

BACKGROUND OF THE INVENTION

[0003] Cyclosporin A (CsA), a neutral cyclic undecapeptide isolated from the
15 fungus *Tolypocladium inflatum* and currently marketed as Neoral[®] and Sandimmune[®] (Novartis, Basel, Switzerland), has been widely used for the prevention of organ transplant rejection. The molecular basis for the immunosuppressant activity of cyclosporin A and cyclosporin analogues begins with the passive diffusion of the cyclosporin (Cs) molecule into the cell, followed by binding to its intracellular
20 receptor, cyclophilin A (CypA). CypA belongs to a family of proteins that catalyze *cis-trans* peptidyl-prolyl isomerization, i.e., PPIase, a rate-limiting step in protein folding. CsA and other cyclosporin analogues bind to the active site of CypA. However, immunosuppression is not believed to be due to the inhibition of CypA PPIase activity. The target of the CsA-CypA complex is a Ca²⁺-calmodulin-
25 dependent serine-threonine-specific protein phosphatase, calcineurin. In T-cells responding to antigen presentation, an increase in intracellular Ca²⁺ activates calcineurin, which subsequently dephosphorylates the transcription factor called the nuclear factor of activated T-cells ("NFAT"). Dephosphorylated NFAT undergoes a molecular change, e.g., homodimerization that allows it to cross into the nucleus, and
30 promotes the expression of T-cell activation genes. CsA and other immunosuppressive cyclosporin derivatives inhibit calcineurin which results in the

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inhibition of expression of cytokine genes, e.g., interleukin-2 (IL-2) that promotes T-cell activation and proliferation, i.e., immunosuppressive activity.

5 *Human Immunodeficiency Viruses and Cyclosporin A or Non-Immunosuppressive Cyclosporins*

[0004] Human immunodeficiency viruses ("HIVs") are lentiviruses, a family of mammalian retroviruses evolved to establish chronic persistent infection with gradual onset of clinical symptoms. There are two major families of HIV. Most of the epidemic involves HIV-1; HIV-2 is a close relative whose distribution is concentrated in western Africa.

[0005] Human cyclophilins A and B have been identified as cellular proteins which bind specifically to HIV-1 Gag polyprotein, p55^{gag}. Gag proteins play a major role in several steps of the virus life cycle, including the assembly and release of virions (Willis et al., "Form, Function, and Use of Retroviral Gag Proteins," *AIDS* 5:639-654 (1991)). A cleavage product of the Gag polyprotein, the capsid protein, has been shown to bind specifically to cyclophilin A. Cyclophilin A is functionally associated with the HIV-1 virions through interaction with the Gag polyprotein. This interaction between cyclophilin A and Gag proteins is inhibited by the immunosuppressive drug, cyclosporin A (Thali et al., "Functional Association of Cyclophilin A With HIV-1 Virions," *Nature* 372:363-365 (1994)).

[0006] Cyclosporin A has demonstrated *in vitro* antiviral activity against HIV-1 (Karpas et al., "Inhibition of Human Immunodeficiency Virus and Growth of Infected T-cells by the Immunosuppressive Drugs Cyclosporin A and FK 506," *Proc. Natl. Acad. Sci. USA* 89:8351-8355 (1992)); however, initial *in vivo* studies in which cyclosporin A was administered as a monotherapy in HIV-infected patients at advanced stages of disease did not show a beneficial effect from the treatment (Levy et al., "Long-Term Follow-Up of HIV Positive Asymptomatic Patients Having Received Cyclosporin A," *Adv. Ex. Med. Biol.* 374:229-234 (1995)). U.S. Patent No. 4,814,323 to Andrieu et al. reported that administration of cyclosporins may be used for the prevention of AIDS in patients infected with the virus before the appearance of the AIDS symptoms, that is patients with no symptoms or patients with AIDS related complex.

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[0007] Highly active antiretroviral therapy (“HAART”) has dramatically decreased the HIV-related morbidity and mortality rates among HIV-infected patients and the transmission of HIV from mother to child by efficiently suppressing viral replication (Palella et al., “Declining Morbidity and Mortality Among Patients With
5 Advanced Human Immunodeficiency Virus Infection,” *N. Eng. J. Med.* 338:853-860 (1998)). Limitations of HAART have become better understood. Thus, the virus can be suppressed to undetectable levels but not eradicated. In addition, there is an ever-growing list of side effects, the eventual development of resistance, and the cost and complexity of HAART regimens that must be contended with.

10 [0008] HAART covers a broad range of antiretroviral agents that include nucleoside reverse transcriptase inhibitors (“NRTI”), nonnucleoside reverse transcriptase inhibitors (“NNRTI”), HIV protease inhibitors, and fusion inhibitors. Specific examples of antiviral agents from each of these families include: Zidovudine, Didanosine, Stavudine, and Lamivudine from the NRTI antiviral class; Nevirapine,
15 Efavirenz, and Delavirdine from the NNRTI antiviral class; Saquinovir, Indinavir, and Ritonavir from the HIV protease inhibitor class; and Enfuvirtide from the fusion inhibitor antiviral class.

[0009] From an immunological standpoint, the introduction of HAART allows for only a partial immune reconstitution. Indeed, *ex vivo* measures of immune
20 function do not generally normalize and, most importantly, HIV-specific T cell responses remain almost invariably impaired. Though several variables have been identified that correlate with the degree of immune reconstitution during HAART, the actual underlying mechanism(s) responsible for such an incomplete immune reconstitution are still poorly understood and likely reflect the severe HIV-driven
25 perturbations in T cell dynamics and homeostasis and the interaction between host and viral factors (Douek, “Disrupting T-Cell Homeostasis: How HIV-1 Infection Causes Disease,” *AIDS Rev.* 5:172-177 (2003)).

[0010] A strategy aimed at the broadest immune reconstitution, possibly overcoming the limitations of HAART, consists in the adjuvant use of
30 immunomodulants. By combining cyclosporin A with HAART, the goal is to contain the immune activation, either virus-specific or owing to non-specific “by-stander” activation. Results from pilot studies in HIV-infected patients has shown that the rapid shutdown of T-cell activation induced by cyclosporin A has produced a more

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rapid and stable increase in CD4+ T-cells and a significant long-term increase in IFN- γ secreting CD4+ and CD4+CCR7- T-cells, establishing a more favorable immunological set-point (Bandera et al., "Immunomodulants in HIV Infection," *Expert Opin. Ther. Patents* 15(9):1115-1131 (2005)). Determination of the long-term efficacy must be assessed in order to understand if this approach truly has value.

[0011] SDZ NIM 811 is a cyclosporin analogue that is completely devoid of immunosuppressive activity but exhibits potent and selective anti-HIV-1 activity (Mlynar et al., "The Non-Immunosuppressive Cyclosporin A Analogue SDZ NIM 811 Inhibits Cyclophilin A Incorporation Into Virions and Virus Replication in Human Immunodeficiency Virus Type-1-Infected Primary and Growth-Arrested Cells," *J. General Virology* 78:825-835 (1997)). SDZ NIM 811 does not prevent the activation of CD4+ T-cell activation as cyclosporin A does. In a manner similar to cyclosporin A, it is proposed that SDZ NIM 811 interferes with the HIV-1 Gag-cyclophilin A interaction to effect its antiviral activity.

[0012] SDZ NIM 811 does not inhibit calcineurin and possesses none of the immunosuppressive activity of cyclosporin A. The potent inhibition of calcineurin by cyclosporin, in addition to being responsible for the potent immunosuppressive activity of cyclosporin A, is also believed to be the cause of the toxicity and the narrow therapeutic index of this drug. Separation of immunosuppressive and antiviral activity could lead to novel antiviral cyclosporins with fewer side effects and improved therapeutic index. Elucidation of structure activity relationships for cyclosporins permits the design of non-immunosuppressive cyclosporin derivatives that retain potent (cyclophilin A) PPIase activity to achieve this goal (Bartz et al., "Inhibition of Human Immunodeficiency Virus Replication by Non-Immunosuppressive Analogs of Cyclosporin A," *Proc. Natl. Acad. Sci. USA* 92:5381-5385 (1995)). European Patent No. 484 281, U.S. Patent No. 5,767,069, U.S. Patent No. 5,948,884, and French Patent Nos. 2,757,520, 2,757,521, and 2,757,522 disclose non-immunosuppressive cyclosporins with antiviral activity.

Hepatitis C Virus and Cyclosporin A

[0013] Recently, cyclosporin A, the most widely prescribed immunosuppressive drug, was reported to be clinically effective against hepatitis C

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viral (HCV) infection (Nakagawa et al., "Specific Inhibition of Hepatitis C Virus Replication by Cyclosporin A," *Biochem. Biophys. Res. Commun.* 313:42-47 (2004)). The authors of the Nakagawa et al. paper state that certain chaperone activities, such as those of cyclophilins, may be crucial for the processing and maturation of the viral proteins and for viral replication.

[0014] A subsequent controlled clinical trial showed that a combination of cyclosporin A with interferon $\alpha 2b$ is more effective than interferon monotherapy, especially in patients with high viral loads (Inoue et al., "Combined Interferon $\alpha 2b$ and Cyclosporin A in the Treatment of Chronic Hepatitis C: Controlled Trial," *J. Gastroenterol.* 38:567-572 (2003)).

[0015] PCT International Patent Publication No. WO 2006/005610 recently described the use of a combination of cyclosporin A and pegylated interferon for treating hepatitis C viral infection. In addition, PCT International Patent Publication No. WO 2005/021028 relates to the use of non-immunosuppressive cyclosporins for treatment of HCV disorders. Also, Paeshuyse et al., "Potent and Selective Inhibition of Hepatitis C Virus Replication by the Non-Immunosuppressive Cyclosporin Analogue DEBIO-025," *Antiviral Research* 65(3):A41 (2005) recently published results for a non-immunosuppressive cyclosporin analogue, DEBIO-025, that exhibited potent and selective inhibition of hepatitis C virus replication. Notably, the cyclosporin derivative DEBIO-025 is also effective for the treatment of HIV-1 (Rosenwirth et al., "Debio-025, A Novel Non-Immunosuppressive Cyclosporine Analog with Potent Anti-Human Immunodeficiency Virus Type 1 Activity: Pharmacological Properties and Mode of Action," *Antiviral Research* 65(3):A42-A43 (2005)). Debio-025 does possess potent binding affinity for cyclophilin A.

[0016] There is still a large need for novel cyclosporin analogues that have therapeutic utility in the treatment of viral-induced diseases.

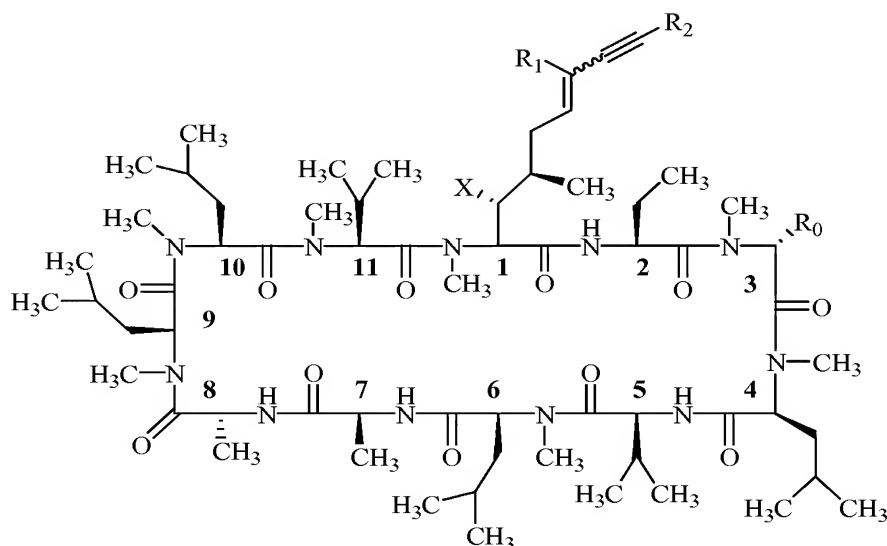
[0017] The present invention is directed to achieving these objectives.

SUMMARY OF THE INVENTION

[0018] The present invention relates to a method of preventing or treating a mammal with a viral-induced disorder. The method involves administering to the

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mammal a therapeutically effective amount of a compound having the following formula:



Formula I

where:

10 X is OH or OAc;

R₀ is H, CH₂OH, or CH₂OR₃;

R₁ is hydrogen, deuterium, or methyl;

15 R₂ is selected from the group consisting of:

hydrogen;

halogen;

C₁-C₆ saturated or unsaturated, straight or branched carbon chain;

20 C₁-C₆ saturated or unsaturated, straight or branched carbon chain containing a substitution or substitutions selected from the group consisting of deuterium, halogen, nitrogen, sulfur, and silicon;

25 C₁-C₆ saturated or unsaturated, straight or branched carbon chain containing a function group or function groups selected from the group consisting of alcohol, ether, aldehyde, ketone, carboxylic acid, ester, and amide;

C₁-C₆ saturated or unsaturated, straight or branched carbon chain containing a function group of oxime or hydrazone;

C₁-C₆ saturated or unsaturated, straight or branched carbon chain containing an aryl or a heteroaryl group;

30 C₃-C₆ substituted and unsubstituted cycloalkyl;
substituted and unsubstituted aryl; and
substituted and unsubstituted heteroaryl; and

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R₃ is selected from the group consisting of:

5 alkanoyl,
 alkenoyl,
 alkynoyl,
 aryloyl,
 arylalkanoyl,
 alkylaminocarbonyl,
 arylaminocarbonyl,
10 arylalkylaminocarbonyl,
 alkyloxy carbonyl,
 aryloxy carbonyl, and
 arylalkyloxy carbonyl,

15 where the compound is a cis geometric isomer, a trans geometric isomer, or a mixture of the cis and the trans geometric isomers or a pharmaceutically acceptable salt thereof,

under conditions effective to prevent or treat the viral-induced disorder.

20 **[0019]** The present invention discloses chemically modified cyclosporin analogues containing a carbon-carbon triple bond on the side chain of the position one amino acid and optionally a substitution on the position three amino acid of cyclosporin A. In particular, the present invention discloses novel cyclosporin analogues containing a conjugated system of a carbon-carbon triple bond and a
25 carbon-carbon double bond on the side chain of the position one amino acid of cyclosporin A.

[0020] The present invention discloses novel cyclosporin analogues which are effective as antiviral agents. The cyclosporin derivatives of the present invention used to treat viral infections may possess potent immunosuppressive activity (via
30 inhibition of calcineurin) or may be completely devoid of immunosuppressive activity (do not inhibit calcineurin). However, the mechanism that the immunosuppressive and non-immunosuppressive cyclosporin compounds share is their activity at cyclophilin A.

35 **BRIEF DESCRIPTION OF THE DRAWING**

[0021] Figure 1 depicts the results from a concanavalin A (ConA)-stimulated murine splenocyte assay, where the novel cyclosporin analogue compounds of the

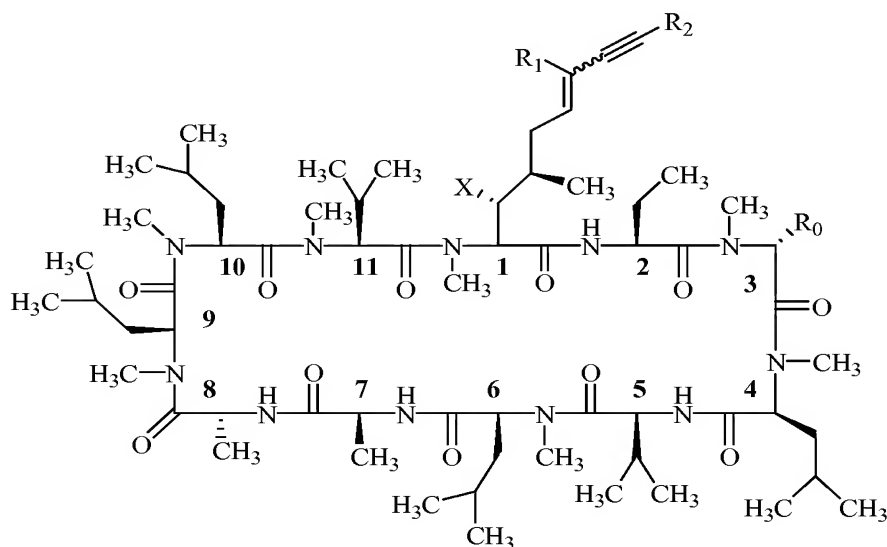
present invention (disclosed in Examples 5 and 33) are shown to possess enhanced or similar potency in immunosuppression, compared to cyclosporin A and ISA_{TX}247.

DETAILED DESCRIPTION OF THE INVENTION

5

[0022] The present invention relates to a method of preventing or treating a mammal with a viral-induced disorder. The method involves administering to the mammal a therapeutically effective amount of a compound having the following formula:

10



Formula I

15 where:

X is OH or OAc;

R₀ is H, CH₂OH, or CH₂OR₃;

20

R₁ is hydrogen, deuterium, or methyl;

R₂ is selected from the group consisting of:

hydrogen;

25 halogen;

C₁-C₆ saturated or unsaturated, straight or branched carbon chain;

C₁-C₆ saturated or unsaturated, straight or branched carbon chain containing a substitution or substitutions selected from the group consisting of deuterium, halogen, nitrogen, sulfur, and silicon;

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C₁-C₆ saturated or unsaturated, straight or branched carbon chain containing a function group or function groups selected from the group consisting of alcohol, ether, aldehyde, ketone, carboxylic acid, ester, and amide;

5 C₁-C₆ saturated or unsaturated, straight or branched carbon chain containing a function group of oxime or hydrazone;

C₁-C₆ saturated or unsaturated, straight or branched carbon chain containing an aryl or a heteroaryl group;

10 C₃-C₆ substituted and unsubstituted cycloalkyl;
substituted and unsubstituted aryl; and
substituted and unsubstituted heteroaryl; and

R₃ is selected from the group consisting of:

15 alkanoyl,
alkenoyl,
alkynoyl,
aryloyl,
arylalkanoyl,
alkylaminocarbonyl,
arylamino carbonyl,
20 arylalkylaminocarbonyl,
alkyloxy carbonyl,
aryloxy carbonyl, and
arylalkyloxy carbonyl,

25 where the compound is a cis geometric isomer, a trans geometric isomer, or a mixture of the cis and the trans geometric isomers or a pharmaceutically acceptable salt thereof,

under conditions effective to prevent or treat the viral-induced disorder.

30 [0023] The wavy bond in the structure of Formula I indicates the carbon-carbon double bond is the cis geometric isomer, the trans geometric isomer, or a mixture of both the cis and the trans geometric isomers.

[0024] One embodiment of the present invention is the above compound of Formula I, where: X is OH or OAc; R₀ is H, CH₂OH, or CH₂OAc; R₁ is H or D; and
35 R₂ is H.

[0025] Another embodiment of the present invention is the above compound of Formula I, where: X is OH or OAc; R₀ is H, CH₂OH, or CH₂OAc; R₁ is H or D; and R₂ is CH₃, CD₃, CH₂CH₃, or CH₂CH₂CH₃.

[0026] Another embodiment of the present invention is the above compound
40 of Formula I, where: X is OH or OAc; R₀ is H, CH₂OH, or CH₂OAc; R₁ is H or D; and R₂ is -CH=CH₂, -CH=CHCH₃, -C≡CH, or -C≡C-CH₃.

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[0027] Another embodiment of the present invention is the above compound of Formula I, where: X is OH or OAc; R₀ is H, CH₂OH, or CH₂OAc; R₁ is H or D; and R₂ is F, Cl, Br, or I.

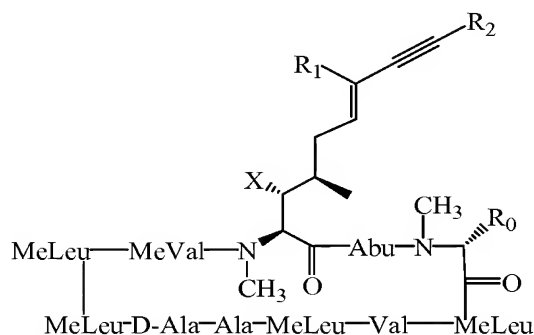
[0028] Another embodiment of the present invention is the above compound of Formula I, where: X is OH or OAc; R₀ is H, CH₂OH, or CH₂OAc; R₁ is H or D; and R₂ is cyclopropyl.

[0029] Another embodiment of the present invention is the above compound of Formula I, where: X is OH or OAc; R₀ is H, CH₂OH, or CH₂OAc; R₁ is H or D; and R₂ is CH₂OH, CH(OH)CH₃, CH₂OCH₃, CH₂OCH₂CH₃, CHO, and C(=O)CH₃.

[0030] Another embodiment of the present invention is the above compound of Formula I, where: X is OH or OAc; R₀ is H, CH₂OH, or CH₂OAc; R₁ is H or D; and R₂ is CH=N-OCH₃, CH=N-OCH₂CH₃, CH=N-NHCH₃, or CH=N-N(CH₃)₂.

[0031] Another embodiment of the present invention is the above compound of Formula I, where: X is OH or OAc; R₀ is H; R₁ is H; and R₂ is H, CH₃, Si(CH₃)₃.

[0032] Other embodiments of the present invention include novel cyclosporine analogue compounds represented by Formula Ia, as shown below:



Formula Ia

20 where:

X = OH or OAc;

R₀ = H;

25

R₁ = H or D; and

R₂ = H, CH₃, CD₃, n-propyl, n-butyl, -CH₂CH₂CH₂Cl, Si(CH₃)₃, Ph, *p*-ClC₆H₄-, 2-pyridinyl, 3-thiophenyl, cyclopropyl, cyclohexyl, -CH₂OH, -CH(OH)CH₃,

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-CH₂CH₂OH, -CH(OH)CH₂CH₃, -CHO, -C(=O)CH₃, -CH=CH₂, -CH=CHCH₃,
 -C(CH₃)=CH₂, -CH=CHI, Br, I, -CH₂OCH₃, -CH₂OPh, -CH₂OCH₂CH=CH₂,
 -CH₂OCH₂C≡CH, -CH₂SCH₂CH₃, -CH₂OCH₂CO₂Bu^t, -CH₂OCH₂CO₂H,
 -CO₂H, -CH₂CH₂CO₂H, -CH=N-OCH₃, or -CH=N-N(CH₃)₂;

5

or where:

X = OH or OAc;

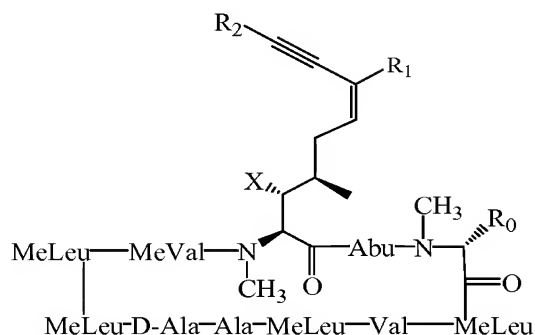
10 R₀ = CH₂OH or CH₂OAc;

R₁ = H or D; and

15 R₂ = H, CH₃, CD₃, n-propyl, Si(CH₃)₃, Br, -CH=CH₂, 3-thiophenyl, -CH₂OH,
 -CH(OH)CH₃, -CH₂OCH₃, -CH₂SCH₂CH₃, -CH₂OPh, -CH₂OCH₂C≡CH,
 cyclopropyl, or cyclohexyl.

[0033] Other embodiments of the present invention include novel
 cyclosporine analogue compounds represented by Formula Ib, as shown below:

20



Formula Ib

where:

25 X = OH or OAc;

R₀ = H;

R₁ = H or D; and

30

R₂ = H, CH₃, Si(CH₃)₃.

[0034] In particular, the present invention relates to novel cyclosporin
 analogues containing a carbon-carbon triple bond on the side chain of the position one
 35 amino acid and optionally a substitution on the position three amino acid of

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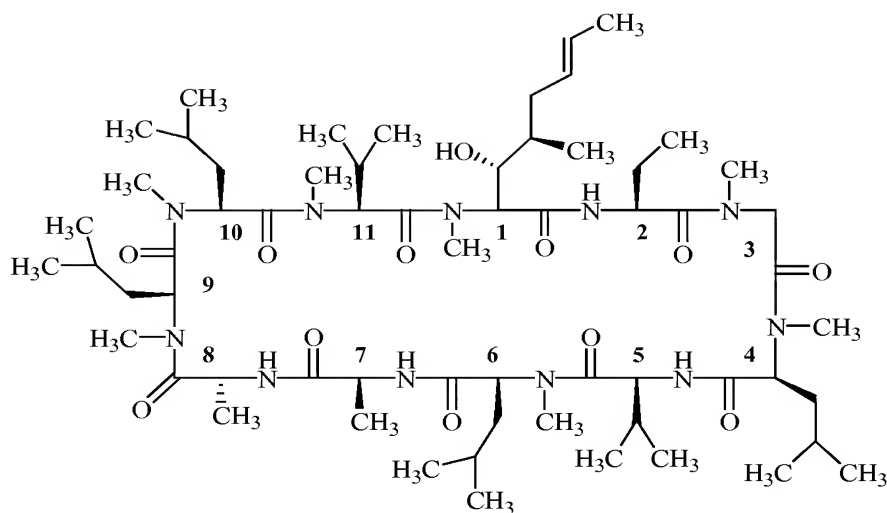
cyclosporin A. More particularly, the present invention relates to novel cyclosporin analogues containing a conjugated system, in which a function group of a carbon-carbon triple bond conjugating with a carbon-carbon double bond is incorporated.

[0035] A carbon-carbon triple bond exists in many natural products (Gung et al., "Total synthesis of (S)-(-)-(E)-15,16-Dihydrominquartynoic Acid: A Highly Potent Anticancer Agent," *J. Org. Chem.*, 69:3488-3492 (2004); Ito et al., "Cytotoxic Polyacetylenes from the Twigs of *Ochanostachys amentacea*," *J. Nat. Prod.*, 64:246-248 (2001), which are hereby incorporated by reference in their entirety). It is well known to use alkynes as pharmaceutical agents. However, only one cyclosporin alkyne, in which a carbon-carbon triple bond replaces the carbon-carbon double bond on the side chain of the position one amino acid of cyclosporin A, is known in the literature. Unfortunately, this modification significantly reduces the immunosuppressive activity of cyclosporin A, where this known cyclosporin alkyne shows only 10% relative immunosuppressive activity compared to cyclosporin A (Rich et al., "Synthesis, Conformation, and Immunosuppressive Activities of Three Analogues of Cyclosporin A Modified in the 1-Position," *J. Med. Chem.*, 33:999-1009 (1990), which is hereby incorporated by reference in its entirety). In contrast, the novel cyclosporin alkyne analogues of the present invention, which contains a conjugated system of a carbon-carbon triple bond and a carbon-carbon double bond, possess enhanced immunosuppressive activity over cyclosporin A.

[0036] The present invention also discloses methods for preparing the compounds represented by Formula I.

[0037] The starting material for the preparation of compounds of the present invention is cyclosporin A. The structure of cyclosporin A, a cycloundecapeptide, and the position numbering for each amino acid in the ring is shown below:

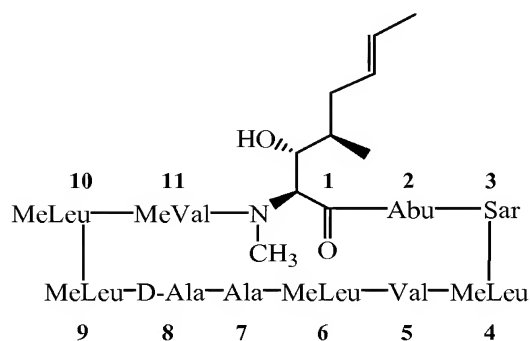
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Cyclosporin A (CsA)

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[0038] Cyclosporin A can also be represented by Formula IIa, as shown below:



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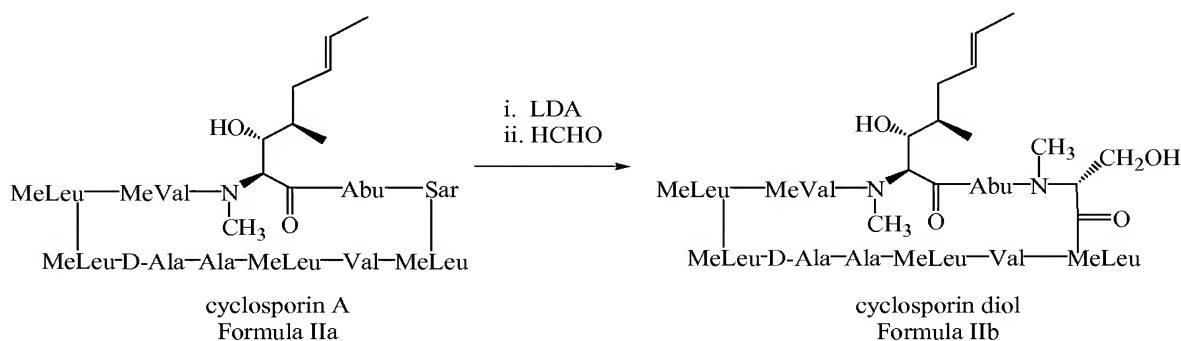
Formula IIa

[0039] The novel cyclosporin analogues of the present invention are derived from cyclosporin A or cyclosporin diol (Formula IIb), a key intermediate prepared by modification at the position three amino acid of cyclosporin A. As shown in Scheme 1, the cyclosporin diol intermediate can be prepared by deprotonation of cyclosporin A with lithium diisopropylamide (LDA) followed by treatment with formaldehyde (Seebach et al, "Modification of Cyclosporin A: Generation of an Enolate at the

15

Sarcosine Residue and Reaction with Electrophiles,” *Helv. Chim. Acta*, 76:1564-1590 (1993), which is hereby incorporated by reference in its entirety).

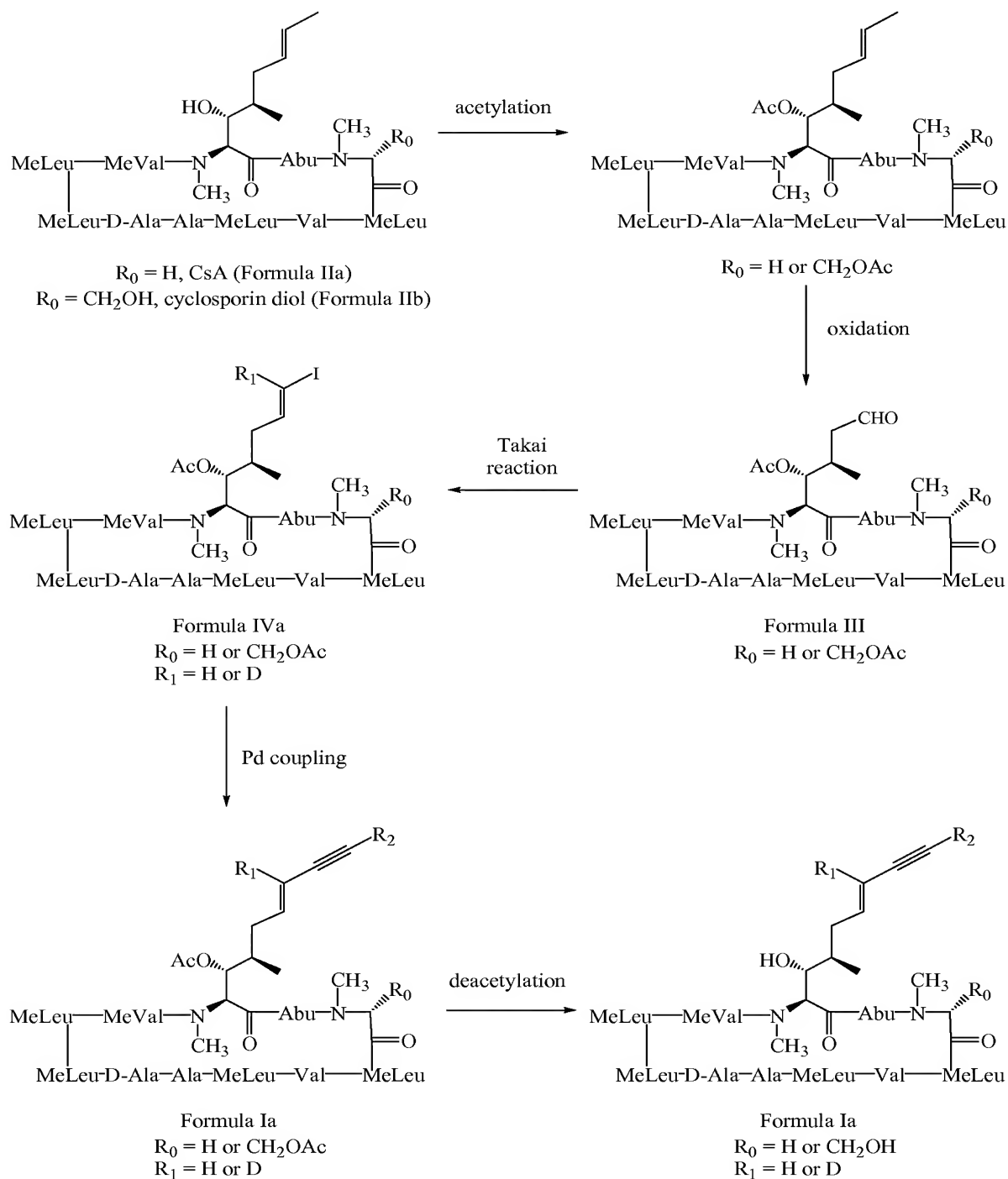
Scheme 1



[0040] According to one embodiment of the present invention, novel cyclosporin alkyne analogues can be prepared by employing palladium-mediated couplings (such as Sonogashira and Negishi coupling) as a key step, which is outlined in Scheme 2. Acetylation of cyclosporin A (Formula IIa) or cyclosporin diol intermediate of Formula IIb with acetic anhydride, followed by oxidative cleavage of the double bond with ozone, generates the cyclosporin aldehyde of Formula III smoothly. Treatment of the cyclosporin aldehyde with iodoform-CrCl₂ complex (Takai et al, “Simple and Selective Method for RCHO → (E)-RCH=CHX Conversion by Means of a CHX₃-CrCl₂ System,” *J. Am. Chem. Soc.*, 108:7408-7410 (1986), which is hereby incorporated by reference in its entirety) affords cyclosporin vinyl iodides of Formula IVa in good to excellent yield (50-80%) in exclusively the trans-configuration. Palladium-catalyzed coupling (such as Sonogashira coupling and Negishi coupling) of the cyclosporin vinyl iodide of Formula IVa with various alkynes or alkynylzinc reagents provides novel cyclosporin alkynes of Formula Ia in the trans-configuration in high yield. The acetyl protecting group in the cyclosporin alkyne of Formula Ia can be removed by treatment with potassium carbonate in methanol (see Scheme 2).

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Scheme 2



- 5 [0041] Utilizing the same strategy, the cis-isomer of cyclosporin alkyne analogues (Formula 1b) can be prepared using the cis-isomer of cyclosporin vinyl iodide (Formula IVb), as shown in Scheme 3. By application of phosphorous ylide chemistry (Wittig reaction), the cyclosporin aldehyde of Formula III can be converted

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to the cyclosporin vinyl iodide of Formula IVb exclusively as the cis-isomer.

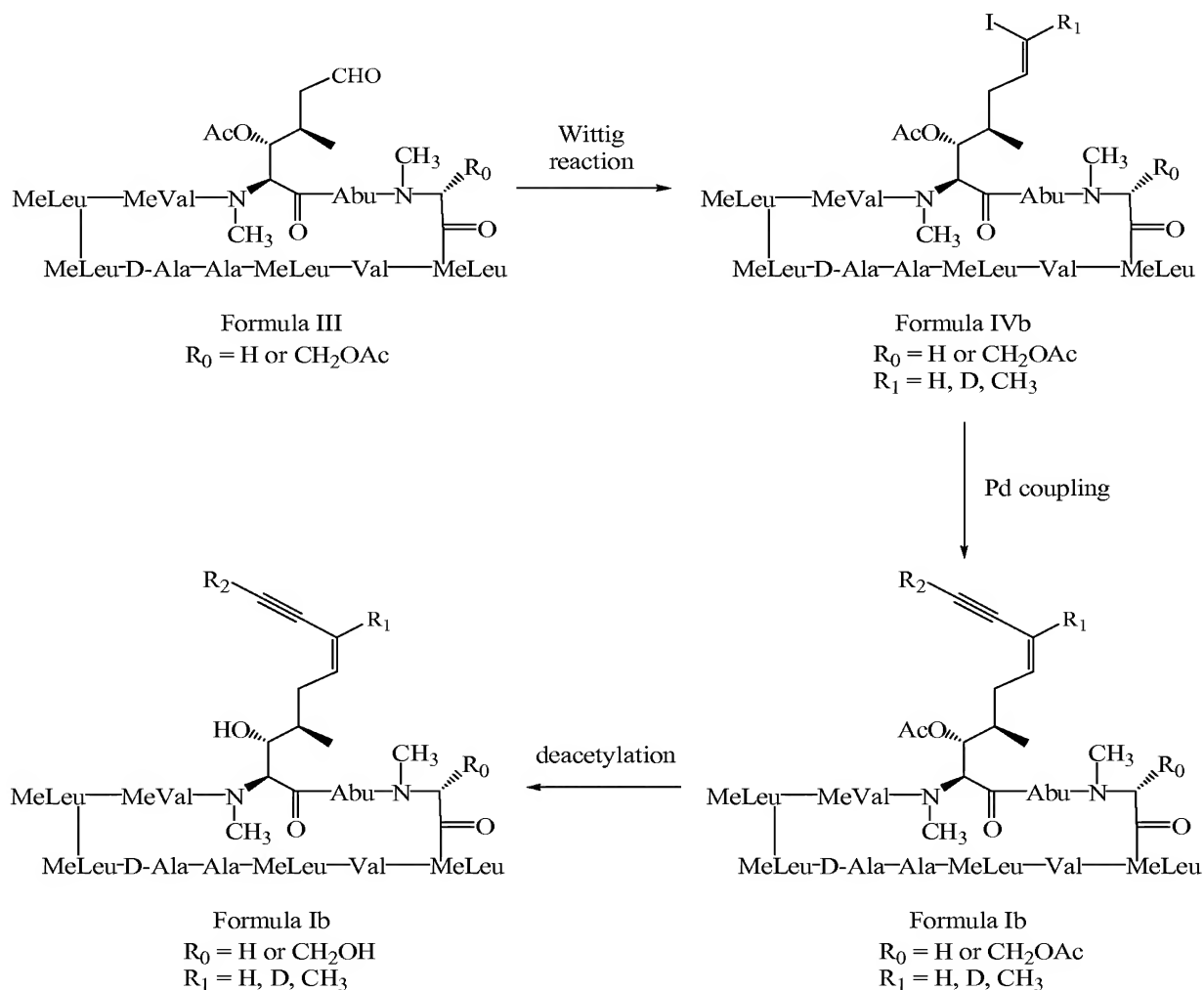
Typically, the phosphorous ylide can be generated by treatment of the corresponding phosphonium salt with a strong base, such as *n*-butyllithium or sodium

bis(trimethylsilyl)amide. Palladium catalyzed coupling (such as Sonogashira

- 5 coupling and Negishi coupling) of the cyclosporin vinyl iodide of Formula IVb with various alkynes or alkynylzinc reagents provides the novel cyclosporin alkyne of Formula Ib in the cis-configuration. The deacetylation is conducted under the same conditions as described above in Scheme 2.

10

Scheme 3



[0042] According to another embodiment of the present invention,

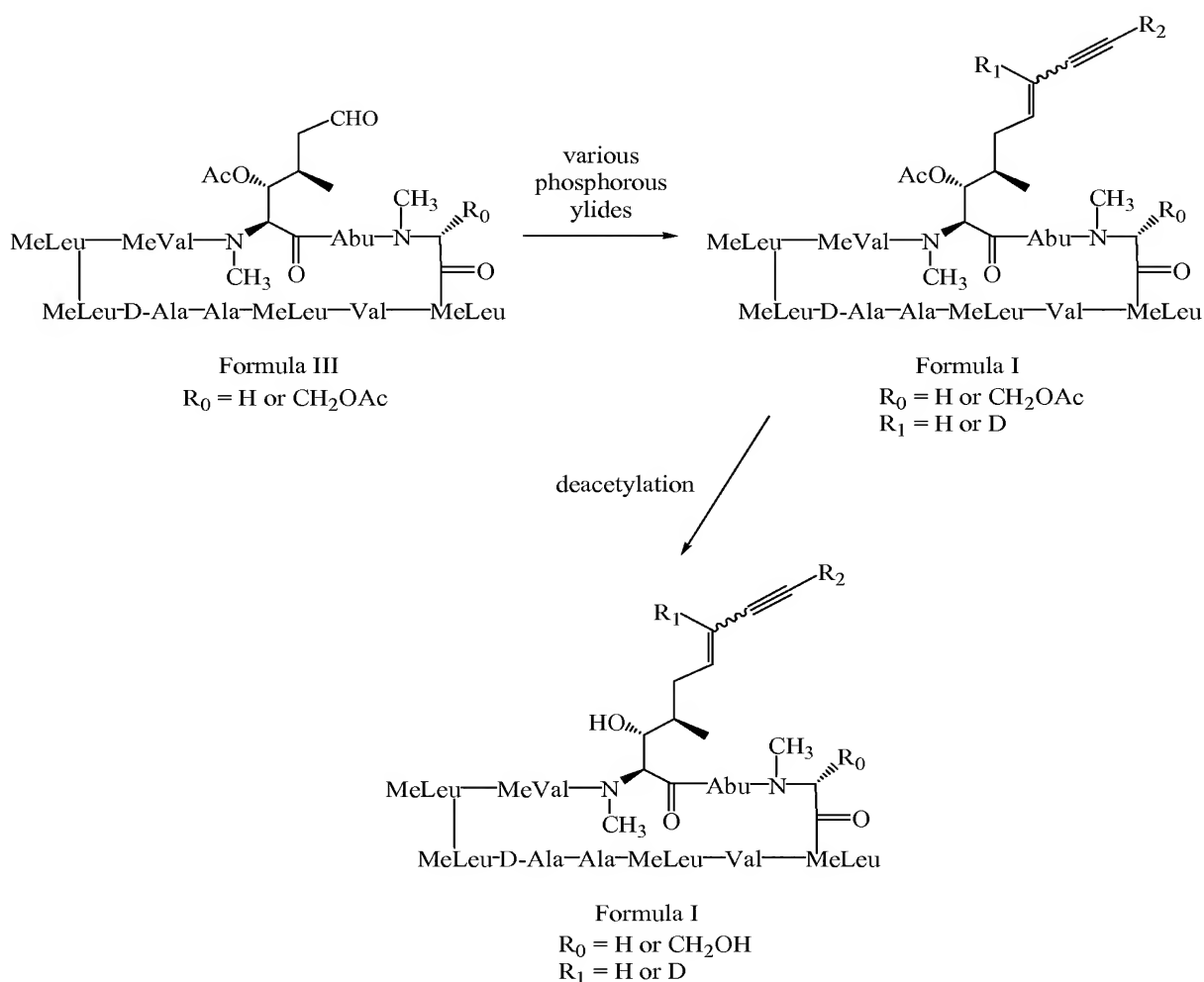
cyclosporin alkyne analogues can be prepared by the application of phosphorus

- 15 chemistry (such as Wittig reaction), as shown in Scheme 4. This chemistry converts

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the cyclosporin aldehyde of Formula III to a mixture of cis and trans geometric isomers of cyclosporin alkyne analogues of Formula I, which is difficult to be separated. Typically, the phosphorous ylide species under Wittig or Horner-Emmons conditions are generated by treatment of various phosphonium salts or phosphonates with a strong base, such as *n*-butyllithium or sodium bis(trimethylsilyl)amide. The deacetylation is conducted under the same conditions as described in Scheme 2.

Scheme 4



10

[0043] Some of the compounds disclosed in the present invention are useful as immunosuppressive agent. Administration of these compounds suppresses the immune response in organ transplant patients and, thus, prevents allograft rejection.

15 The compounds of the present invention possess enhanced immunosuppressive

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activity over cyclosporin A. For example, as shown in Figure 1, the novel cyclosporin alkyne analogue compound disclosed in Example 5 demonstrates immunosuppressive activity up to two times more potent (as determined by IC_{50}) over ISA_{TX}247 and up to three or four times more active, compared to cyclosporin A, while the cyclosporin alkyne analogue compound disclosed in Example 33 is up to two times more active over cyclosporin A and equipotent to ISA_{TX}247 in the concanavalin A (ConA) stimulated murine splenocyte assay. Table 1 shows the immunosuppressive activities of several novel cyclosporin alkyne analogue compounds disclosed in the present application. (The third column in Table 1 contains Cyclosporin A positive control values included for comparison.)

Table 1. Immunosuppressive Activity of Novel Cyclosporin Alkyne Analogue Compounds of the Present Invention

Example where the Novel Cyclosporin Alkyne Analogue Compound is Disclosed	IC_{50} (ng/mL)	IC_{50} (ng/mL) of CsA
Example 5	6	20
Example 6	6	23
Example 13	1	15
Example 15	8	17
Example 27	6	9
Example 28	7	5
Example 33	9	20
Example 71	4	15
Example 73	6	15
Example 79	11	15
Example 83	10	15
Example 89	15	17

15

[0044] The compounds disclosed in the present invention are useful for the prevention or treatment of viral-induced disorders that are dependent upon the presence of cyclophilin A. The compounds of the present invention used to treat these viral infections may possess potent immunosuppressive activity (via inhibition of calcineurin) or may be completely devoid of immunosuppressive activity (do not inhibit calcineurin). However, the mechanism that the immunosuppressive and non-immunosuppressive cyclosporin compounds share is their activity at cyclophilin A.

20

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[0045] Cyclophilin A enzyme activity, i.e., peptidyl-prolyl cis-trans isomerase activity, is important to the folding and trafficking of proteins. The HIV infectivity of CD4+ T-cells and viral replication are dependent upon the incorporation of cyclophilin A into HIV-1 virions through interactions with the Gag polyprotein.

5 Inhibition of the cyclophilin A enzyme activity is necessary and sufficient for anti-HIV-1 activity.

[0046] In one embodiment of the present invention, the viral-induced disorder is a human immunodeficiency virus (HIV)-induced disorder. Thus, compounds of the present invention that lack immunosuppressant activity as determined by the
10 Concanavalin A (Con A)-stimulated murine splenocyte assay but retain potent peptidyl prolyl isomerase (PPIase) inhibitory (cyclophilin A) activity may possess anti-HIV activity. In addition, compounds of the present invention that have immunosuppressive activity as determined by the Con A-stimulated murine splenocyte assay and also possess potent PPIase inhibitory (cyclophilin A) activity
15 may possess anti-HIV activity.

[0047] *In vitro* biological assays that allow the determination of binding affinity to cyclophilin A or allow the determination of inhibition of peptidyl cis-trans isomerase activity are described in Handschumacher et al., "Cyclophilin: A Specific Cytosolic Binding Protein for Cyclosporin A," *Science* 226:544-547 (1984) and
20 Kofron et al., "Determination of Kinetic Constants for Peptidyl Prolyl Cis-Trans Isomerases by an Improved Spectrophotometric Assay," *Biochemistry* 30:6127-6134 (1991), respectively, which are both hereby incorporated by reference in their entirety.

[0048] The *in vitro* anti-HIV activity of compounds of the present invention can be measured in established cell line cultures as described by Mayaux et al.,
25 "Triterpene Derivatives That Block Entry of Human Immunodeficiency Virus Type 1 Into Cells," *Proc. Natl. Acad. Sci. USA* 91:3564-3568 (1994), which is hereby incorporated by reference in its entirety.

[0049] In another embodiment of the present invention, the compound of the present invention is administered in combination with antiretroviral agents, such as
30 nucleoside reverse transcriptase inhibitors, nonnucleoside reverse transcriptase inhibitors, human immunodeficiency virus protease inhibitors, fusion inhibitors, and combinations thereof. Examples of nucleoside reverse transcriptase inhibitors include, but are not limited to, Zidovudine, Didanosine, Stavudine, and Lamivudine.

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Examples of nonnucleoside reverse transcriptase inhibitors include, but are not limited to, Nevirapine, Efavirenz, and Delavirdine. Examples of human immunodeficiency virus protease inhibitors include, but are not limited to, Saquinovir, Indinavir, and Ritonavir. Examples of fusion inhibitors include, but are not limited to, Enfuvirtide.

[0050] Although cyclophilin PPIase activity would appear to be implicated in anti-HCV activity as it is for anti-HIV-1 activity, the hepatitis C virus (HCV) proteins that may interact with cyclophilin A have yet to be identified. In another embodiment of the present invention, the viral-induced disorder is a HCV-induced disorder.

10 Hepatitis C infections or HCV induced disorders are, for example, chronic hepatitis, liver cirrhosis, or liver cancer (e.g., hepatocellular carcinoma). Thus, compounds of the present invention that lack immunosuppressant activity as determined by the Concanavalin A (Con A)-stimulated murine splenocyte assay but retain potent peptidyl prolyl isomerase (PPIase) inhibitory (cyclophilin A) activity may possess
15 anti-HCV activity. In addition, compounds of the present invention that have immunosuppressive activity as determined by the Con A-stimulated murine splenocyte assay and also possess potent PPIase inhibitory (cyclophilin A) activity may possess anti-HCV activity. The compounds of the present invention may also be used as a prophylactic treatment for neonates born to HCV-infected mothers, for
20 healthcare workers exposed to the virus, or for transplant recipients, e.g., organ or tissue transplant (e.g. liver transplant) recipients, to eliminate possible recurrent infection after transplantation.

[0051] In another embodiment of the present invention, the compound of the present invention is administered in combination with an interferon. Examples of
25 interferons include, but are not limited to, interferon α 2a and interferon α 2b. The interferon can be a pegylated interferon. Examples of interferons include, but are not limited to, pegylated interferon α 2a or pegylated interferon α 2b.

[0052] Utility of the immunosuppressive or non-immunosuppressive cyclosporin compounds of the present invention in treating diseases or conditions
30 from HCV infection can be demonstrated in standard animal or clinical tests in accordance with the methods described in Examples 102 and 103, for example.

[0053] Some of the compounds disclosed in the present invention also possess utility in the treatment of autoimmune and chronic inflammatory diseases such as

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asthma, rheumatoid arthritis, multiple sclerosis, psoriasis, and ulcerative colitis, to name only a few.

[0054] The compounds disclosed in the present invention are also useful for the treatment of ocular allergy and dry eye. Allergan is currently marketing a topical
5 formulation of cyclosporin A called Restasis™ (cyclosporin ophthalmic emulsion) for the treatment of keratoconjunctivitis sicca or chronic dry eye syndrome in patients whose tear production is presumed to be suppressed due to ocular inflammation. While the exact mechanism of Restasis™ is unknown, it is thought to act as an immunomodulator with anti-inflammatory effects (“Annual Update 2003: Ophthalmic
10 Drugs” *Drugs of the Future*, 28(3): 287-307 (2003); Clark et al., “Ophthalmic Drug Discovery,” *Nature Reviews in Drug Discovery*, 2:448-459 (2003), which are hereby incorporated by reference in their entirety).

[0055] For treatment of the above mentioned diseases, therapeutically effective doses of the compounds of the present invention may be administered orally,
15 topically, parenterally, by inhalation spray, or rectally in dosage unit formulations containing conventional non-toxic pharmaceutically acceptable carriers, adjuvants, and vehicles. The term parenteral, as used herein, includes subcutaneous injections, intravenous, intramuscular, intrasternal injection, or infusion techniques.

[0056] The pharmaceutical compositions containing the active ingredient may
20 be in the form suitable for oral use, for example, as tablets, troches, lozenges, aqueous or oily suspensions, dispersible powders or granules, emulsions, hard or soft capsules, or syrups or elixirs. The pharmaceutical compositions of the present invention contain the active ingredient formulated with one or more pharmaceutically acceptable carriers. As used herein, the term “pharmaceutical acceptable carrier”
25 means a non-toxic, inert solid, semi-solid or liquid filler, diluent, encapsulating material, or formulation auxiliary of any type. Some examples of pharmaceutically acceptable carriers are sugars such as lactose, glucose, and sucrose; starches such as corn starch or potato starch; cellulose and its derivatives such as sodium carboxymethyl cellulose, ethyl cellulose, and cellulose acetate; powdered tragacanth;
30 malt; gelatin; talc; excipients such as cocoa butter and suppository waxes; oils such as peanut oil, cottonseed oil, safflower oil, sesame oil, olive oil, corn oil, and soybean oil; glycols such as propylene glycol; esters such as ethyl oleate and ethyl laurate; agar; buffering agents such as magnesium hydroxide and aluminum hydroxide;

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alginic acid; pyrogen-free water; isotonic saline; Ringer's solution; ethyl alcohol; phosphate buffer solutions; non-toxic, compatible lubricants such as sodium lauryl sulfate and magnesium stearate; as well as coloring agents, releasing agents, sweetening, and flavoring and perfuming agents. Preservatives and antioxidants, such as ethyl or n-propyl p-hydroxybenzoate, can also be included in the pharmaceutical compositions.

[0057] Dosage forms for topical or transdermal administration of compounds disclosed in the present invention include ointments, pastes, creams, lotions, gels, plasters, cataplasms, powders, solutions, sprays, inhalants, or patches. The active component is admixed under sterile conditions with a pharmaceutically acceptable carrier and any needed preservatives or buffers, as may be required. The ointments, pastes, creams and gels may contain, in addition to an active compound of the present invention, excipients such as animal and vegetable fats, oils, waxes, paraffins, starch, tragacanth, cellulose derivatives, polyethylene glycols, silicones, bentonites, silicic acid, talc and zinc oxide, or mixtures thereof.

[0058] For nasal administration, the compounds disclosed in the present invention can be administered, as suitable, in liquid or powdered form from a nasal applicator. Forms suitable for ophthalmic use will include lotions, tinctures, gels, ointment and ophthalmic inserts, as known in the art. For rectal administration (topical therapy of the colon), the compounds of the present invention may be administered in suppository or enema form, in solution in particular, for example in vegetable oil or in an oily system for use as a retention enema.

[0059] Compounds disclosed in the present invention may be delivered to the lungs by the inhaled route either in nebulizer form or as a dry powder. The advantage of the inhaled route, over the systemic route, in the treatment of asthma and other diseases of airflow obstruction and/or chronic sinusitis, is that patients are exposed to very small quantities of the drug and the compound is delivered directly to the site of action.

[0060] Dosages of the compounds of the present invention employed for the treatment of the maladies identified in the present invention will vary depending on the site of treatment, the particular condition to be treated, the severity of the condition, the subject to be treated (who may vary in body weight, age, general health, sex, and other factors) as well as the effect desired.

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[0061] Dosage levels ranging from about 0.05 mg to about 50 mg per kilogram of body weight per day are useful for the treatment of the conditions or diseases identified in the present invention. This means the amount of the compound disclosed in the present invention that is administered will range from 2.5 mg to about 2.5 gm per patient per day.

[0062] The amount of active ingredient that may be combined with the pharmaceutical carrier materials to produce a single dosage form will vary depending upon the host treated and the particular mode of administration. For example, a formulation intended for the oral administration of humans may contain from 2.5 mg to 2.5 gm of active compound of the present invention compounded with an appropriate and convenient amount of carrier material which may vary from about 5 to 95 percent of the total composition. Dosage unit forms will generally contain between from about 5 mg to about 500 mg of active compound of the present invention. Dosage for topical preparation will, in general be less (one tenth to one hundredth) of the dose required for an oral preparation.

EXAMPLES

Example 1 – Preparation of Cyclosporin Acetate

[0063] A solution of cyclosporin A (5.0 g, 4.16 mmol), acetic anhydride (7.80 mL, 83.2 mmol), and DMAP (760 mg, 6.2 mmol) in methylene chloride (40 mL) was stirred overnight at room temperature under N₂ atmosphere. Saturated sodium bicarbonate solution (200 mL) was added to the solution and stirred for an additional 2 h. The mixture was extracted with ether, washed with 1 N HCl, neutralized with saturated sodium bicarbonate solution, washed with brine, dried over sodium sulfate, and concentrated *in vacuo* to afford cyclosporin acetate (4.92 g, 95%) as a white solid: ¹H NMR (300 MHz, CDCl₃) δ 8.57 (d, *J* = 9.6 Hz, 1H), 8.04 (d, *J* = 6.9 Hz, 1H), 7.51 (d, *J* = 9.4 Hz, 1H), 7.47 (d, *J* = 7.8 Hz, 1H), 5.67 (dd, *J* = 11.0, 4.0 Hz, 1H), 5.60–5.44 (m, 2H), 5.39 (dd, *J* = 11.7, 3.7 Hz, 1H), 5.32–5.13 (m, 4H), 5.06–4.93 (m, 2H), 4.85 (t, *J* = 7.2 Hz, 1H), 4.77 (t, *J* = 9.6 Hz, 1H), 4.65 (d, *J* = 13.7 Hz, 1H), 4.41 (t, *J* = 7.0 Hz, 1H), 3.46 (s, 3H), 3.26 (s, 3H), 3.24 (s, 3H), 3.21 (s, 3H), 3.10 (s, 3H), 2.68 (s, 3H), 2.66 (s, 3H), 2.50–2.35 (m, 1H), 2.25–1.80 (m, 6H),

2.08 (s, 3H), 2.01 (s, 3H), 1.75–1.55 (m, 6H), 1.45–0.75 (m, 55H); ESI MS m/z 1245 $[\text{C}_{64}\text{H}_{113}\text{N}_{11}\text{O}_{13} + \text{H}]^+$.

Example 2 – Preparation of Acetyl Cyclosporin Aldehyde

5 **[0064]** Ozone was bubbled into a solution of cyclosporin acetate from Example 1 (3.0 g, 2.4 mmol) in methylene chloride (70 mL) at -78°C until a blue color was developed. The mixture was degassed with nitrogen for a few min and dimethylsulfide (3 mL) was added at -78°C . The reaction mixture was allowed to
10 warm to room temperature and stirred for 3 h. The reaction mixture was concentrated *in vacuo* and the residue was dissolved in ethyl acetate (300 mL), washed with water (2×70 mL) and brine (70 mL), dried over sodium sulfate, filtered, and concentrated *in vacuo* to afford acetyl cyclosporin aldehyde (2.79 g, 94%) as a white solid. The crude was carried to the next step without further purification: ^1H NMR (300 MHz, CDCl_3) δ 9.60 (d, $J = 3.5$ Hz, 1H), 8.55 (d, $J = 9.7$ Hz, 1H), 7.96 (d, $J = 6.8$ Hz, 1H),
15 7.52 (d, $J = 7.7$ Hz, 1H), 7.46 (d, $J = 9.0$ Hz, 1H), 5.67 (dd, $J = 11.0, 3.8$ Hz, 1H), 5.60–5.45 (m, 2H), 5.32 (dd, $J = 12.1, 3.3$ Hz, 1H), 5.24–5.10 (m, 2H), 5.08–4.90 (m, 2H), 4.84 (t, $J = 7.1$ Hz, 1H), 4.73 (t, $J = 9.6$ Hz, 1H), 4.64 (d, $J = 13.8$ Hz, 1H), 4.41 (t, $J = 7.0$ Hz, 1H), 3.46 (s, 3H), 3.29 (s, 6H), 3.21 (s, 3H), 3.08 (s, 3H), 2.67 (s, 3H),
20 2.65 (s, 3H), 2.50–2.35 (m, 2H), 2.25–1.80 (m, 6H), 1.99 (s, 3H), 1.75–1.55 (m, 3H), 1.50–0.75 (m, 57H); ESI MS m/z 1233 $[\text{C}_{62}\text{H}_{109}\text{N}_{11}\text{O}_{14} + \text{H}]^+$.

Example 3 – Preparation of trans Acetyl Cyclosporin Vinyl Iodide

25 **[0065]** To an ice-cooled suspension of chromium(II) chloride (1.0 g, 8.2 mmol) in THF (25 mL) was added a solution of acetyl cyclosporin aldehyde from Example 2 (0.50 g, 0.41 mmol) and iodoform (1.29 g, 3.28 mmol) in THF (25 mL). After 7 h at 0°C , the reaction mixture was poured into ice-water (50 mL). The water layer was extracted with ethyl acetate (3×60 mL). The combined organics were
30 dried over anhydrous sodium sulfate and concentrated. The material was purified by semi-preparative HPLC to afford trans acetyl cyclosporin vinyl iodide (290 mg, 52%) as a white solid: ^1H NMR (300 MHz, CDCl_3) δ 8.47 (d, $J = 9.8$ Hz, 1H), 8.01 (d, $J = 6.4$ Hz, 1H), 7.56 (d, $J = 8.8$ Hz, 1H), 7.54 (d, $J = 7.5$ Hz, 1H), 6.50–6.40 (m, 1H), 5.84 (d, $J = 14.3$ Hz, 1H), 5.69–5.10 (m, 6H), 4.97 (d, $J = 11.1$ Hz, 2H), 4.87–4.73

(m, 2H), 4.64 (d, $J = 13.8$ Hz, 1H), 4.43 (t, $J = 7.0$ Hz, 1H), 3.43 (s, 3H), 3.28 (s, 3H), 3.26 (s, 3H), 3.20 (s, 3H), 3.12 (s, 3H), 2.68 (s, 3H), 2.66 (s, 3H), 2.45–2.35 (m, 1H), 2.28–1.80 (m, 8H), 2.06 (s, 3H), 1.77–1.60 (m, 3H), 1.50–0.75 (m, 56H); ESI MS m/z 1357 $[\text{C}_{63}\text{H}_{110}\text{IN}_{11}\text{O}_{13} + \text{H}]^+$.

5

Example 4 – Preparation of the Acetate of Cyclosporin Alkyne

[0066] A mixture of acetyl cyclosporin vinyl iodide from Example 3 (90 mg, 0.066 mmol), CuI (6 mg, 0.033 mmol), and $\text{PdCl}_2(\text{PPh}_3)_2$ (23 mg, 0.033 mmol) in
10 Et_3N (3 mL) was stirred at room temperature under N_2 atmosphere. Reaction mixture was then treated with (trimethylsilyl)acetylene (65 mg, 0.66 mmol). Reaction was allowed to stir for 3 h at room temperature. Solution was filtered through a micro-filter and then concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford the acetate of cyclosporin alkyne (37 mg, 42%) as an off-
15 white solid: ^1H NMR (300 MHz, CDCl_3) δ 8.47 (d, $J = 9.6$ Hz, 1H), 8.03 (d, $J = 6.9$ Hz, 1H), 7.52 (d, $J = 9.0$ Hz, 1H), 7.47 (d, $J = 7.7$ Hz, 1H), 6.12–6.04 (m, 2H), 5.68 (dd, $J = 10.9, 3.7$ Hz, 2H), 5.53 (s, 1H), 5.48 (s, 1H), 5.36–5.11 (m, 7H), 5.01–4.62 (m, 10H), 4.41 (t, $J = 7.0$ Hz, 2H), 4.06 (t, $J = 6.6$ Hz, 2H), 3.44 (s, 3H), 3.27 (s, 3H), 3.26 (s, 3H), 3.20 (s, 3H), 3.07 (s, 3H), 2.68 (s, 2H), 2.65 (s, 3H), 2.04 (s, 2H),
20 2.00 (s, 3H), 1.31 (d, $J = 7.1$ Hz, 2H), 1.26 (d, $J = 7.0$ Hz, 2H), 1.06–0.79 (m, 50H), 0.17 (s, 9H); ESI MS m/z 1327 $[\text{C}_{68}\text{H}_{119}\text{N}_{11}\text{O}_{13}\text{Si} + \text{H}]^+$.

Example 5 - Preparation of Cyclosporin Alkyne

25 [0067] A solution of the acetate of cyclosporin alkyne from Example 4 (37 mg, 0.02 mmol) in methanol (2 mL) was stirred at room temperature. Reaction mixture was treated with potassium carbonate (30 mg, 0.21 mmol) and allowed to keep stirring under N_2 atmosphere overnight. Mixture was diluted with ethyl acetate, washed with saturated sodium bicarbonate solution and brine, dried over sodium
30 sulfate, and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford the cyclosporin alkyne (15 mg, 45%) as a white solid: ^1H NMR (300 MHz, CDCl_3) δ 7.94 (d, $J = 9.8$ Hz, 1H), 7.64 (d, $J = 7.4$ Hz, 1H), 7.41 (d, $J = 8.3$ Hz, 1H), 7.16 (d, $J = 7.8$ Hz, 1H), 6.31–6.20 (m, 2H), 5.70 (dd, $J = 10.8, 3.7$ Hz, 2H), 5.50 (d, $J = 5.9$ Hz, 2H), 5.42 (s, 1H), 5.36 (s, 1H), 5.32 (d, $J = 3.7$ Hz,

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- 1H), 5.28 (d, $J = 3.9$ Hz, 1H), 5.11–4.92 (m, 6H), 4.84 (t, $J = 7.1$ Hz, 2H), 4.47 (s, 1H), 4.75–4.62 (m, 4H), 4.13 (s, 1H), 3.91–3.80 (m, 3H), 3.51 (s, 3H), 3.39 (s, 3H), 3.26 (s, 3H), 3.10 (s, 6H), 2.70 (s, 3H), 2.68 (s, 3H), 1.34 (d, $J = 7.2$ Hz, 2H), 1.29–1.24 (m, 8H), 1.08–0.81 (m, 45H), 0.74 (d, $J = 6.4$ Hz, 2H); ESI MS m/z 1213
- 5 [C₆₃H₁₀₉N₁₁O₁₂ + H]⁺; HPLC >99% (AUC), $t_R = 13.58$ min.

Example 6 - Preparation of Cyclosporin Alkyne

- [0068] To a solution of 1-propynylmagnesium bromide (7.3 mL, 0.5 M in THF, 3.6 mmol) under stirring and nitrogen was added zinc chloride (3.6 mL, 1 M in ether, 3.6 mmol). After 30 min at room temperature, a solution of acetyl cyclosporin vinyl iodide from Example 3 (500 mg, 0.36 mmol) in THF (5 mL) was added, followed by dichlorobis(triphenylphosphine)palladium(II) (258 mg, 0.36 mmol). After 18 h at room temperature, the solvent was removed under reduced pressure.
- 15 The crude material was purified by semi-preparative HPLC to afford the acetate of cyclosporin alkyne (335 mg, 71%) as a pale yellow solid: ESI MS m/z 1269 [C₆₆H₁₁₃N₁₁O₁₃ + H]⁺.
- [0069] To a stirred solution of the above cyclosporin alkyne (335 mg, 0.26 mmol) in methanol (10 mL) was added potassium carbonate (360 mg, 2.6 mmol) at room temperature. After 18 h, ethyl acetate (100 mL) and water (50 mL) were added. The organic layer was separated, dried over anhydrous sodium sulfate, and concentrated under vacuum to afford crude product. The material was purified by semi-preparative HPLC to afford cyclosporin alkyne (53 mg, 16%) as an off-white solid: ¹H NMR (300 MHz, CDCl₃) δ 7.89 (d, $J = 9.5$ Hz, 1H), 7.70 (d, $J = 7.2$ Hz, 1H), 7.50 (d, $J = 8.2$ Hz, 1H), 7.26 (d, $J = 5.2$ Hz, 1H), 6.03–5.93 (m, 1H), 5.70 (dd, $J_1 = 4.0$ Hz, $J_2 = 11.0$ Hz, 1H), 5.50 (d, $J = 5.8$ Hz, 1H), 5.40–5.29 (m, 3H), 5.16–4.65 (m, 8H), 4.49 (t, $J = 7.0$ Hz, 1H), 3.82–3.78 (m, 1H), 3.49 (s, 3H), 3.36 (s, 3H), 3.23 (s, 3H), 3.11 (s, 3H), 3.09 (s, 3H), 2.72 (s, 3H), 2.70 (s, 3H), 2.60–0.70 (m, 70H); ESI MS m/z 1227 [C₆₄H₁₁₁N₁₁O₁₂ + H]⁺; HPLC >99% (AUC), $t_R = 14.17$ min.
- 25
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Example 7 - Preparation of 1-(Trimethylsilyl)propyne-*d*₃

- [0070] Trimethylsilylacetylene (4.9 mL, 34.5 mmol) is dissolved in THF (20 mL) and cooled to – 78°C. *n*-BuLi (15.2 mL, 2.5 M in hexanes, 38 mmol) was

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added to the reaction dropwise and stirred for 0.5 h at -78°C . Iodomethane- d_3 (5.0 g, 38 mmol) was added to the reaction dropwise. Resulting mixture was warmed to room temperature over 1 h. Reaction was quenched with water and extracted with ethyl ether (3×100 mL). Combined organics were washed with water, brine, dried
5 over sodium sulfate and distilled using a short path distillation setup to remove most of solvents. Residue was transferred to a small flask and distilled to give 1-(trimethylsilylpropyne- d_3 (1.8 g, 32%) as colorless oil: ^1H NMR (300 MHz, CDCl_3) δ 0.13 (s, 9H); ESI MS m/z 116 [$\text{C}_6\text{H}_9\text{D}_3\text{Si} + \text{H}$] $^+$.

10 **Example 8 – Preparation of Deuterated Cyclosporin Alkyne**

[0071] 1-(Trimethylsilyl)propyne- d_3 from Example 7 (64 mg, 0.5mmol) was dissolved in Et_3N (3 mL) and cooled to 0°C . Solution was treated with Bu_4NF (0.66 mL, 1 M in THF, 0.66 mmol) and allowed to stir at 0°C under N_2 atmosphere
15 for 10 min. Reaction was allowed to warm to room temperature. After warming, the reaction mixture was treated with acetyl cyclosporin vinyl iodide from Example 3 (75 mg, 0.05 mmol), CuI (5 mg, 0.02 mmol), and $\text{PdCl}_2(\text{PPh}_3)_2$ (19 mg, 0.02 mmol). Reaction was kept stirring for 3 h at room temperature. Solution was filtered through a micro-filter and then concentrated *in vacuo*.

20 [0072] The above crude product (32 mg, 0.025 mmol) was dissolved in methanol (2 mL) and stirred at room temperature. Reaction mixture was treated with potassium carbonate (35 mg, 0.25 mmol) and allowed to keep stirring under N_2 atmosphere overnight. Mixture was diluted with ethyl acetate, washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated *in*
25 *vacuo*. The crude product was purified by semi-preparative HPLC to afford deuterated cyclosporin alkyne (15 mg, 50%) as a white solid: ^1H NMR (300 MHz, CDCl_3) δ 7.98 (d, $J = 11.5$ Hz, 1H), 7.62 (d, $J = 7.5$ Hz, 1H), 7.42 (d, $J = 8.3$ Hz, 1H), 7.15 (d, $J = 7.9$ Hz, 1H), 6.06–5.96 (m, 2H), 5.70 (dd, $J = 10.8, 3.8$ Hz, 2H), 5.50 (d, $J = 5.9$ Hz, 2H), 5.39 (s, 1H), 5.35–5.28 (m, 2H), 5.13–4.90 (m, 10H), 4.82 (t, $J =$
30 7.2 Hz, 2H), 4.77 (s, 1H), 4.75–4.62 (m, 6H), 4.50 (t, $J = 7.0$ Hz, 2H), 3.51 (s, 3H), 3.39 (s, 3H), 3.26 (s, 3H), 3.10 (s, 6H), 2.70 (s, 3H), 2.68 (s, 3H), 1.34 (d, $J = 7.2$ Hz, 2H), 1.25 (t, $J = 3.0$ Hz, 4H), 1.09–0.81 (m, 45H), 0.71 (d, $J = 6.1$ Hz, 2H); ESI MS m/z 1230 [$\text{C}_{64}\text{H}_{108}\text{D}_3\text{N}_{11}\text{O}_{12} + \text{H}$] $^+$; HPLC >99% (AUC), $t_R = 13.74$ min.

Example 9 - Preparation of *trans*-Cyclosporin Vinyl Iodide

[0073] To a stirred solution of *trans*-acetyl cyclosporin vinyl iodide from
5 Example 3 (200 mg, 0.15 mmol) in methanol (20 mL) was added potassium carbonate
(518 mg, 3.75 mmol) at room temperature. After 12 h at room temperature, methanol
was evaporated. Water (30 mL) was added and the mixture was extracted with ethyl
acetate (3 × 80 mL). The organic layer was separated, dried over anhydrous sodium
sulfate, and concentrated under vacuum to afford crude product. The material was
10 purified by semi-preparative HPLC to afford *trans*-cyclosporin vinyl iodide (125 mg,
63%) as a white solid: ¹H NMR (300 MHz, CDCl₃) δ 7.90 (d, *J* = 9.3 Hz, 1H), 7.66
(d, *J* = 6.5 Hz, 1H), 7.49 (d, *J* = 8.5 Hz, 1H), 7.18 (d, *J* = 7.8 Hz, 1H), 6.55–6.43 (m,
1H), 5.93 (d, *J* = 14.0 Hz, 1H), 5.69 (d, *J* = 8.1 Hz, 1H), 5.47 (d, *J* = 5.9 Hz, 1H), 5.32
(d, *J* = 8.0 Hz, 1H), 5.12–4.92 (m, 4H), 4.82 (t, *J* = 6.2 Hz, 1H), 4.74 (d, *J* = 14.8 Hz,
15 1H), 4.67 (t, *J* = 9.1 Hz, 1H), 4.53 (t, *J* = 7.2 Hz, 1H), 3.82 (t, *J* = 6.2 Hz, 1H), 3.50 (s,
3H), 3.37 (s, 3H), 3.25 (s, 3H), 3.11 (s, 6H), 2.72 (s, 3H), 2.69 (s, 3H), 2.48–1.90 (m,
8H), 1.80–1.53 (m, 6H), 1.50–0.72 (m, 55H); ESI MS *m/z* 1315 [C₆₁H₁₀₈IN₁₁O₁₂ +
H]⁺; HPLC >99% (AUC), *t*_R = 14.35 min.

Example 10 - Preparation of Cyclosporin Alkyne

[0074] A mixture of cyclosporin vinyl iodide from Example 9 (82 mg,
0.062 mmol), CuI (6 mg, 0.031 mmol), and PdCl₂(PPh₃)₂ (22 mg, 0.031 mmol) in
Et₃N (3 mL) was stirred at room temperature under N₂ atmosphere. Reaction mixture
25 was then treated with (trimethylsilyl)acetylene (61 mg, 0.62 mmol). Reaction was
allowed to stir for 3 h at room temperature. Solution was filtered through a micro-
filter and then concentrated *in vacuo*. The crude product was purified by semi-
preparative HPLC to afford cyclosporin alkyne (30 mg, 37%) as an off-white solid:
¹H NMR (300 MHz, CDCl₃) δ 7.89 (d, *J* = 9.7 Hz, 1H), 7.61 (d, *J* = 7.5 Hz, 1H), 7.48
30 (d, *J* = 8.0 Hz, 1H), 7.13 (d, *J* = 7.8 Hz, 1H), 6.21–6.10 (m, 2H), 5.71 (dd, *J* = 10.5,
3.6 Hz, 2H), 5.53 (d, *J* = 5.3 Hz, 2H), 5.47 (s, 1H), 5.42 (s, 1H), 5.27 (dd, *J* = 11.4,
3.8 Hz, 2H), 5.21–4.99 (m, 6H), 4.93–4.71 (m, 4H), 4.62 (t, *J* = 9.3 Hz, 2H), 4.55 (d,
J = 7.3 Hz, 2H), 3.51 (s, 3H), 3.38 (s, 3H), 3.26 (s, 3H), 3.10 (s, 3H), 3.09 (s, 3H),
2.70 (s, 3H), 2.69 (s, 3H), 1.34 (d, *J* = 7.2 Hz, 2H), 1.32–1.24 (m, 9H), 1.08–0.83 (m,

45H), 0.68 (d, $J = 5.7$ Hz, 2H), 0.17 (s, 9H); ESI MS m/z 1285 [$C_{66}H_{117}N_{11}O_{12}Si + H$] $^+$; HPLC 97.1% (AUC), $t_R = 22.14$ min.

Example 11 - Preparation of Cyclosporin Alkyne

5 [0075] The acetate of cyclosporin alkyne from Example 4 (488 mg, 0.52 mmol) was dissolved in THF (10 mL) and cooled to 0°C. Solution was treated with Bu₄NF (6.3 mL, 1 M in THF, 6.3 mmol) and allowed to stir for 20 min under N₂ atmosphere. Reaction was quenched with saturated sodium bicarbonate solution and
10 warmed to room temperature. Reaction mixture was extracted with ethyl acetate (3 × 100 mL). Combined organic layers were washed with brine, dried over sodium sulfate and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford cyclosporin alkyne (231 mg, 50%) as an off-white solid:
15 ¹H NMR (300 MHz, CDCl₃) δ 8.51 (d, $J = 9.7$ Hz, 1H), 8.00 (d, $J = 6.7$ Hz, 1H), 7.71–7.63 (m, 2H), 7.56–7.43 (m, 2H), 7.38 (d, $J = 9.1$ Hz, 1H), 6.26–6.15 (m, 2H), 5.67 (dd, $J = 7.1, 3.8$ Hz, 2H), 5.53 (d, $J = 5.9$ Hz, 2H), 5.38 (dd, $J = 11.9, 3.9$ Hz, 2H), 5.31–5.26 (m, 2H), 5.14 (t, $J = 7.6$ Hz, 2H), 4.97 (d, $J = 11.0$ Hz, 2H), 4.94–4.73 (m, 4H), 4.46–4.33 (m, 8H), 3.44 (s, 3H), 3.27 (s, 3H), 3.25 (s, 3H), 3.21 (s, 3H), 3.09 (s, 3H), 2.67 (s, 3H), 2.65 (s, 3H), 2.01 (s, 3H), 1.32–1.23 (m, 4H), 1.06–0.70 (m,
20 50H); ESI MS m/z 1255 [$C_{65}H_{111}N_{11}O_{13} + H$] $^+$.

Example 12 - Preparation of the Acetate of Cyclosporin Alkynyl Iodide

[0076] A mixture of cyclosporin alkyne from Example 11 (65 mg, 0.05 mmol)
25 and CuI (100 mg, 0.52 mmol) in THF (3 mL) was cooled to –15°C. Bis-(trimethylsilyl)peroxide (90 mg, 0.52 mmol) was added to the reaction dropwise under N₂ atmosphere. The reaction was kept stirring at –15°C for 15 min and then allowed to warm to room temperature. Reaction was heated to 50°C and stirred overnight. Reaction mixture was diluted with THF, washed with a saturated solution
30 of ammonium chloride, washed with brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford the acetate of cyclosporin alkynyl iodide (8 mg, 11%) as an off-white solid:
¹H NMR (300 MHz, CDCl₃) δ 8.44 (d, $J = 9.6$ Hz, 1H), 8.02 (d, $J = 6.6$ Hz, 1H), 7.64 (d, $J = 9.3$ Hz, 1H), 7.58 (d, $J = 7.5$ Hz, 1H), 6.09–5.98 (m, 2H), 5.68 (d, $J = 11.1$ Hz,

2H), 5.55–5.06 (m, 10H), 5.97 (d, $J = 10.9$ Hz, 2H), 4.88–4.72 (m, 8H), 4.64 (d, $J = 13.7$ Hz, 2H), 4.44 (t, $J = 6.8$ Hz, 2H), 3.43 (s, 3H), 3.25 (s, 3H), 3.23 (s, 3H), 3.19 (s, 3H), 3.10 (s, 3H), 2.68 (s, 3H), 2.66 (s, 3H), 2.01 (s, 3H), 1.40–1.25 (m, 4H), 1.10–0.71 (m, 50H); ESI MS m/z 1381 [$C_{65}H_{110}IN_{11}O_{13} + H$] $^{+}$.

5

Example 13 - Preparation of Cyclosporin Alkynyl Iodide

[0077] A solution of the acetate of cyclosporin alkynyl iodide from Example 12 (8 mg, 0.006 mmol) in methanol (1 mL) was stirred at room temperature.

10 Reaction mixture was treated with potassium carbonate (20 mg, 0.14 mmol) and allowed to keep stirring under N_2 atmosphere overnight. Mixture was diluted with ethyl acetate, washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford cyclosporin alkynyl iodide (3.2 mg, 46%) as a white

15 solid: 1H NMR (300 MHz, $CDCl_3$) δ 7.95 (d, $J = 9.7$ Hz, 1H), 7.64 (d, $J = 7.4$ Hz, 1H), 7.47 (d, $J = 8.0$ Hz, 1H), 7.21 (d, $J = 7.7$ Hz, 1H), 6.23–6.20 (m, 2H), 5.70 (dd, $J = 10.8, 3.7$ Hz, 2H), 5.55 (s, 1H), 5.48 (d, $J = 6.0$ Hz, 2H), 5.28 (dd, $J = 11.3, 3.7$ Hz, 2H), 5.14–4.91 (m, 6H), 4.82 (t, $J = 7.2$ Hz, 2H), 4.77 (s, 1H), 4.70–4.62 (m, 3H), 4.52 (t, $J = 7.1$ Hz, 2H), 3.81 (d, $J = 6.4$ Hz, 2H), 3.50 (s, 3H), 3.39 (s, 3H), 3.26 (s,

20 3H), 3.10 (s, 6H), 2.69 (s, 3H), 2.68 (s, 3H), 1.34 (d, $J = 7.2$ Hz, 2H), 1.31–1.22 (m, 9H), 1.08–0.80 (m, 45H), 0.72 (d, $J = 6.5$ Hz, 2H); ESI MS m/z 1339 [$C_{63}H_{108}IN_{11}O_{12} + H$] $^{+}$; HPLC 95.4% (AUC), $t_R = 19.54$ min.

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Example 14 - Preparation of the Acetate of Cyclosporin Alkynyl Bromide

[0078] A solution of cyclosporin alkyne from Example 11 (96 mg, 0.08 mmol) and CuBr (328 mg, 2.3 mmol) in THF (4 mL) was cooled to $-15^{\circ}C$. Bis(trimethylsilyl)peroxide (408 mg, 2.3 mmol) was added to the reaction dropwise, under N_2 atmosphere. Reaction mixture was kept stirring at $-15^{\circ}C$ for 20 min, then

30 allowed to warm to room temperature and stirred an additional 5 h. Reaction was quenched with saturated ammonium chloride solution and extracted with ethyl acetate. Organic layer was washed with brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to

afford the acetate of cyclosporin alkynyl bromide (19 mg, 69%) as an off-white solid: ESI MS m/z 1333 [$C_{65}H_{110}BrN_{11}O_{13} + H$]⁺.

Example 15 - Preparation of Cyclosporin Alkynyl Bromide

5 [0079] A solution of the acetate of cyclosporin alkynyl bromide from Example 14 (19 mg, 0.01 mmol) in methanol (2 mL) was stirred at room temperature. Reaction mixture was treated with potassium carbonate (30 mg, 0.22 mmol) and was allowed to keep stirring under N₂ atmosphere overnight. Mixture was diluted with
10 ethyl acetate, washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford cyclosporin alkynyl bromide (8 mg, 43%) as a white solid: ¹H NMR (300 MHz, CDCl₃) δ 7.96 (d, J = 9.7 Hz, 1H), 7.65 (d, J = 7.2 Hz, 1H), 7.43 (d, J = 8.1 Hz, 1H), 7.18 (d, J = 7.8 Hz, 1H), 6.24–6.14 (m, 2H), 5.73–5.68
15 (m, 1H), 5.48 (d, J = 6.2 Hz, 2H), 5.40 (s, 1H), 5.35 (s, 1H), 5.29 (dd, J = 11.7, 3.8 Hz, 2H), 5.07 (d, J = 10.7 Hz, 2H), 5.01–4.93 (m, 3H), 4.85 (t, J = 7.2 Hz, 2H), 4.74 (s, 1H), 4.69 (s, 1H), 4.63 (s, 1H), 4.49 (t, J = 7.2 Hz, 2H), 3.83 (t, J = 6.5 Hz, 2H), 3.50 (s, 3H), 3.39 (s, 3H), 3.26 (s, 3H), 3.10 (s, 6H), 2.70 (s, 3H), 2.68 (s, 3H),
20 2.00 (s, 6H), 1.35 (d, J = 7.1 Hz, 2H), 1.26 (d, J = 4.0 Hz, 3H), 1.08–0.80 (m, 48H), 0.74 (d, J = 6.4 Hz, 1H); ESI MS m/z 1291 [$C_{63}H_{108}BrN_{11}O_{12} + H$]⁺; HPLC 96.9% (AUC), t_R = 20.07 min.

Example 16 – Preparation of the Acetate of Cyclosporin Alkynyl Amine

25 [0080] A mixture of acetyl cyclosporin vinyl iodide from Example 3 (60 mg, 0.044 mmol), CuI (4 mg, 0.022 mmol), and PdCl₂(PPh₃)₂ (15 mg, 0.022 mmol) in Et₃N (3 mL) was stirred at room temperature under N₂ atmosphere. Reaction mixture was then treated with *N*-methylpropargylamine (30 mg, 0.44 mmol). Reaction was allowed to stir for 6 h at room temperature. Solution was filtered through a micro-
30 filter and then concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford the acetate of cyclosporin alkynyl amine (9.5 mg, 17%) as an off-white solid: ¹H NMR (300 MHz, CDCl₃) δ 8.45 (d, J = 9.7 Hz, 1H), 7.99 (d, J = 6.9 Hz, 1H), 7.44 (t, J = 8.0 Hz, 2H), 5.98–5.89 (m, 2H), 5.67 (dd, J = 10.8, 3.6 Hz, 2H), 5.51 (d, J = 3.0 Hz, 2H), 5.40–5.23 (m, 7H), 5.14 (t, J = 7.6 Hz, 2H), 5.00–4.74

(m, 10H), 4.63 (d, $J = 13.8$ Hz, 2H), 4.42 (t, $J = 7.0$ Hz, 2H), 4.10 (s, 1H), 3.44 (s, 3H), 3.26 (s, 3H), 3.24 (s, 3H), 3.20 (s, 3H), 3.08 (s, 3H), 2.67 (s, 3H), 2.65 (s, 3H), 2.57 (s, 2H), 2.02 (s, 3H), 2.01 (s, 2H), 1.30 (t, $J = 6.8$ Hz, 2H), 1.25 (s, 2H), 1.06–0.78 (m, 50H); ESI MS m/z 1298 [$C_{67}H_{116}N_{12}O_{13} + H$]⁺.

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Example 17 - Preparation of Cyclosporin Alkynyl Amine

[0081] A solution of the acetate of cyclosporin alkynyl amine from Example 16 (9.5 mg, 0.007 mmol) in methanol (2 mL) was stirred at room
10 temperature. Reaction mixture was treated with potassium carbonate (11 mg, 0.07 mmol) and allowed to keep stirring under N₂ atmosphere overnight. Mixture was diluted with ethyl acetate, washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford cyclosporin alkynyl amine (4.5 mg,
15 49%) as a white solid: ¹H NMR (300 MHz, CDCl₃) δ 8.00 (d, $J = 10.6$ Hz, 1H), 7.65 (d, $J = 8.5$ Hz, 1H), 7.61 (d, $J = 6.9$ Hz, 1H), 7.21 (d, $J = 7.9$ Hz, 1H), 6.15–6.04 (m, 2H), 5.67 (dd, $J = 11.0, 4.1$ Hz, 2H), 5.54–5.38 (m, 2H), 5.33 (d, $J = 8.1$ Hz, 1H), 5.08–4.63 (m, 6H), 4.85–4.63 (m, 8H), 4.61 (s, 1H), 1.50 (t, $J = 7.2$ Hz, 2H), 4.41 (s, 1H), 4.26 (d, $J = 1.7$ Hz, 2H), 4.14 (d, $J = 1.7$ Hz, 2H), 4.08–3.97 (m, 2H), 3.50 (s,
20 3H), 3.40 (s, 3H), 3.24 (s, 3H), 3.12 (s, 3H), 3.11 (s, 3H), 2.67 (s, 3H), 2.66 (s, 3H), 2.00 (s, 2H), 1.54 (d, $J = 9.6$ Hz, 2H), 1.46 (s, 1H), 1.38–1.25 (m, 6H), 1.05–0.80 (m, 45H), 0.77 (d, $J = 6.5$ Hz, 2H); ESI MS m/z 1256 [$C_{65}H_{114}N_{12}O_{12} + H$]⁺; HPLC 98.8% (AUC), $t_R = 12.28$ min.

Example 18 – Preparation of the Acetate of Cyclosporin Alkynyl Amine

[0082] A mixture of acetyl cyclosporin vinyl iodide from Example 3 (60 mg, 0.044 mmol), CuI (4 mg, 0.022 mmol), and PdCl₂(PPh₃)₂ (15 mg, 0.022 mmol) in Et₃N (3 mL) was stirred at room temperature under N₂ atmosphere. Reaction mixture
30 was then treated with 1-(dimethylamino)-2-propyne (38 mg, 0.44 mmol). Reaction was allowed to stir for 6 h at room temperature. Solution was filtered through a micro-filter and then concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford the acetate of cyclosporin alkynyl amine (33 mg, 56%) as an off-white solid: ¹H NMR (300 MHz, CDCl₃) δ 8.49 (d, $J = 9.6$ Hz, 1H), 8.02 (d, J

= 6.8 Hz, 1H), 7.45 (t, J = 7.7 Hz, 2H), 6.06–5.95 (m, 2H), 5.67 (dd, J = 11.0, 3.8 Hz, 2H), 5.52 (d, J = 6.8 Hz, 2H), 5.36 (s, 1H), 5.31 (s, 1H), 5.14 (t, J = 7.6 Hz, 2H), 5.00–4.94 (m, 9H), 4.86 (t, J = 7.3 Hz, 2H), 4.75 (t, J = 9.5 Hz, 2H), 4.65 (d, J = 13.8 Hz, 2H), 4.41 (t, J = 7.0 Hz, 2H), 4.12 (dd, J = 14.2, 7.1 Hz, 2H), 3.44 (s, 3H),
5 3.26 (s, 3H), 3.25 (s, 3H), 3.21 (s, 3H), 3.07 (s, 3H), 2.67 (s, 2H), 2.65 (s, 3H), 2.32 (s, 6H), 2.04 (s, 2H), 2.01 (s, 3H), 1.31 (d, J = 7.1 Hz, 2H), 1.26 (d, J = 6.8 Hz, 2H), 1.06–0.59 (m, 50H); ESI MS m/z 1312 [$C_{68}H_{118}N_{12}O_{13} + H$]⁺.

Example 19 - Preparation of Cyclosporin Alkynyl Amine

10 [0083] A solution of the acetate of cyclosporin alkynyl amine from Example 18 (33 mg, 0.02 mmol) in methanol (3 mL) was stirred at room temperature. Reaction mixture was treated with potassium carbonate (34 mg, 0.2 mmol) and allowed to keep stirring under N₂ atmosphere overnight. Mixture was diluted with
15 ethyl acetate, washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford cyclosporin alkynyl amine (17 mg, 55%) as a white solid:
¹H NMR (300 MHz, CDCl₃) δ 7.98 (d, J = 9.7 Hz, 1H), 7.64 (d, J = 7.6 Hz, 1H), 7.48 (d, J = 8.1 Hz, 1H), 7.12 (d, J = 7.9 Hz, 1H), 6.21–6.12 (m, 2H), 5.70 (dd, J = 10.7, 3.6 Hz, 2H), 5.43 (d, J = 5.7 Hz, 2H), 5.47 (s, 1H), 5.42 (s, 1H), 5.28 (dd, J = 11.5, 3.8 Hz, 2H), 5.11–4.99 (m, 8H), 4.92 (dd, J = 9.7, 6.0 Hz, 2H), 4.82 (t, J = 7.4 Hz, 2H), 4.75 (s, 2H), 4.70 (s, 2H), 4.63 (t, J = 9.4 Hz, 2H), 4.54 (t, J = 7.3 Hz, 2H), 3.51 (s, 3H), 3.39 (s, 3H), 3.26 (s, 3H), 3.11 (s, 3H), 3.10 (s, 3H), 2.70 (s, 3H), 2.68 (s, 3H), 2.55 (s, 3H), 1.34 (d, J = 7.2 Hz, 2H), 1.30–1.24 (m, 9H), 1.09–0.81 (m, 45H),
20 0.70 (d, J = 6.0 Hz, 2H); ESI MS m/z 1270 [$C_{66}H_{116}N_{12}O_{12} + H$]⁺; HPLC >99% (AUC), t_R = 12.37 min.

Example 20 – Preparation of the Acetate of Cyclosporin Alkynyl Alcohol

30 [0084] A mixture of acetyl cyclosporin vinyl iodide from Example 3 (300 mg, 0.22 mmol), CuI (21 mg, 0.11 mmol), and PdCl₂(PPh₃)₂ (39 mg, 0.055 mmol) in Et₃N (5 mL) was stirred at room temperature under N₂ atmosphere. Reaction mixture was then treated with propargyl alcohol (124 mg, 2.2 mmol). Reaction was allowed to stir for 3 h at room temperature. Solution was filtered through a micro-filter and then

concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford the acetate of cyclosporin alkynyl alcohol (100 mg, 36%) as an off-white solid: ^1H NMR (300 MHz, CDCl_3) δ 8.44 (d, $J = 9.9$ Hz, 1H), 7.96 (d, $J = 6.9$ Hz, 1H), 7.43 (d, $J = 7.9$ Hz, 2H), 5.68 (dd, $J = 10.9, 3.9$ Hz, 2H), 5.57 (d, $J = 10.2$ Hz, 1H), 5.27 (d, $J = 3.3$ Hz, 2H), 5.23 (d, $J = 3.4$ Hz, 2H), 5.00–4.80 (m, 8H), 4.64 (s, 1H), 4.60 (s, 1H), 4.46–4.25 (m, 5H), 4.06 (t, $J = 6.7$ Hz, 2H), 3.43 (s, 3H), 3.25 (s, 3H), 3.20 (s, 3H), 3.10 (s, 3H), 2.67 (s, 3H), 2.65 (s, 3H), 2.04 (s, 4H), 2.03 (s, 5H), 1.30 (t, $J = 7.6$ Hz, 2H), 1.25 (s, 5H), 1.06–0.76 (m, 51H); ESI MS m/z 1285 [$\text{C}_{66}\text{H}_{113}\text{N}_{11}\text{O}_{14} + \text{H}$] $^+$.

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Example 21 - Preparation of Cyclosporin Alkynyl Alcohol

[0085] A solution of acetate of cyclosporin alkynyl alcohol from Example 20 (32 mg, 0.025 mmol) in methanol (2 mL) was stirred at room temperature. Reaction mixture was treated with potassium carbonate (38 mg, 0.27 mmol) and allowed to keep stirring under N_2 atmosphere overnight. The mixture was diluted with ethyl acetate, washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford cyclosporin alkynyl alcohol (21 mg, 68%) as a white solid: ^1H NMR (300 MHz, CDCl_3) δ 7.60 (d, $J = 9.8$ Hz, 1H), 7.40 (d, $J = 9.0$ Hz, 2H), 7.19 (d, $J = 7.7$ Hz, 1H), 6.10–5.99 (m, 1H), 5.66 (dd, $J = 10.6, 3.8$ Hz, 2H), 5.46 (s, 1H), 5.41–5.34 (m, 4H), 5.14–5.02 (m, 6H), 4.89 (d, $J = 4.6$ Hz, 2H), 4.94 (d, $J = 4.8$ Hz, 2H), 4.83–4.66 (m, 4H), 4.44 (t, $J = 7.1$ Hz, 1H), 4.30 (d, $J = 11.9$ Hz, 2H), 4.11–4.02 (m, 4H), 3.50 (s, 3H), 3.39 (s, 3H), 3.23 (s, 3H), 3.10 (s, 3H), 3.09 (s, 3H), 2.68 (s, 6H), 2.18–2.11 (m, 7H) 1.38 (d, $J = 7.2$ Hz, 5H), 1.24 (d, $J = 6.6$ Hz, 6H), 1.06–0.78 (m, 39H); ESI MS m/z 1243 [$\text{C}_{64}\text{H}_{111}\text{N}_{11}\text{O}_{13} + \text{H}$] $^+$; HPLC >99% (AUC), $t_R = 13.16$ min.

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Example 22 - Preparation of Cyclosporin Alkynyl Aldehyde

[0086] A mixture of the acetate of cyclosporin alkynyl alcohol from Example 20 (100 mg, 0.77 mmol), *N*-methylmorpholine *N*-oxide (14 mg, 0.121 mmol), 4Å molecular sieves (200 mg) was dissolved in methylene chloride (2 mL). Mixture was treated with tetrapropylammonium perruthenate (3 mg,

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0.008 mmol) and stirred at room temperature under N₂ atmosphere. Mixture was kept stirring for 2 h and then filtered through a pad of silica gel. The filtrate was concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford cyclosporin alkynyl aldehyde (88 mg, 89%) as an off-white solid: ¹H NMR (300 MHz, CDCl₃) δ 9.32 (s, 1H), 8.46 (d, *J* = 9.5 Hz, 1H), 8.01 (d, *J* = 6.6 Hz, 1H), 7.68 (d, *J* = 8.9 Hz, 1H), 7.58 (d, *J* = 7.5 Hz, 1H), 6.52–6.41 (m, 2H), 5.68 (dd, *J* = 10.8, 3.5 Hz, 2H), 5.53 (d, *J* = 12.7 Hz, 2H), 5.40–5.13 (m, 15H), 4.96 (d, *J* = 11.0 Hz, 2H), 4.64 (d, *J* = 13.8 Hz, 2H), 4.44 (t, *J* = 6.9 Hz, 2H), 3.98 (s, 2H), 3.44 (s, 3H), 3.26 (s, 2H), 3.24 (s, 3H), 3.20 (s, 3H), 3.11 (s, 3H), 2.67 (s, 3H), 2.66 (s, 3H), 2.02 (s, 3H), 1.33–1.19 (m, 4H), 1.06–0.75 (m, 50H); ESI MS *m/z* 1283 [C₆₆H₁₁₁N₁₁O₁₄ + H]⁺.

Example 23 - Preparation of the Acetate of *trans*-Cyclosporin Vinyl Iodide

[0087] A solution of cyclosporin alkynyl aldehyde from Example 22 (87 mg, 0.068 mmol) and iodoform (232 mg, 0.68 mmol) in THF (3 mL) was cooled to -78°C. After cooling, dry Cr(II)Cl₂ (165 mg, 1.35 mmol) was added to the reaction. The reaction mixture was allowed to warm to 0°C and stirred under N₂ atmosphere for 4 h. Mixture was poured into ice water and extracted with ethyl acetate (3 × 100 mL). The combined organic layers were washed with brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford the acetate of *trans*-cyclosporin vinyl iodide (18 mg, 19%) as an off-white solid: ¹H NMR (300 MHz, CDCl₃) δ 8.38 (d, *J* = 8.9 Hz, 1H), 8.01 (d, *J* = 6.3 Hz, 1H), 7.76 (d, *J* = 8.8 Hz, 1H), 7.63 (d, *J* = 6.7 Hz, 1H), 6.92–6.71 (m, 7H), 6.05–5.94 (m, 2H), 5.52 (d, *J* = 8.0 Hz, 2H), 5.37 (d, *J* = 13.2 Hz, 2H), 5.01–4.55 (m, 10H), 4.47 (t, *J* = 6.9 Hz, 2H), 3.41 (s, 2H), 3.25 (s, 3H), 3.21 (s, 3H), 3.19 (s, 3H), 3.10 (s, 3H), 2.83 (dd, *J* = 10.6, 7.6 Hz, 2H), 2.70 (s, 2H), 2.68 (s, 3H), 2.03 (s, 2H), 2.02 (s, 3H), 1.34–1.25 (m, 7H), 1.06–0.74 (m, 50H); ESI MS *m/z* 1407 [C₆₇H₁₁₂IN₁₁O₁₃ + H]⁺.

Example 24 - Preparation of Cyclosporin Vinyl Iodide

[0088] A solution of the acetate of cyclosporin vinyl iodide from Example 23 (18 mg, 0.012 mmol) in methanol (1 mL) was stirred at room temperature. Reaction mixture was treated with potassium carbonate (50 mg, 0.36 mmol) and allowed to

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keep stirring under N₂ atmosphere overnight. Mixture was diluted with ethyl acetate, washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-

5 ¹H NMR (300 MHz, CDCl₃) δ 7.91 (d, *J* = 9.7 Hz, 1H), 7.64 (d, *J* = 7.2 Hz, 1H), 7.46 (d, *J* = 8.2 Hz, 1H), 7.21 (d, *J* = 7.8 Hz, 1H), 6.79 (s, 1H), 6.74 (s, 2H), 6.65 (d, *J* = 2.0 Hz, 2H), 6.60 (d, *J* = 2.1 Hz, 1H), 6.22–6.11 (m, 2H), 5.79 (dd, *J* = 10.7, 3.5 Hz, 2H), 5.49–5.26 (m, 3H), 5.11–4.62 (m, 6H), 4.49 (t, *J* = 7.1 Hz, 2H), 4.40 (s, 1H), 4.30–4.20 (m, 2H), 4.14 (d, *J* = 1.7 Hz, 2H), 3.50 (s, 3H), 3.38 (s, 3H), 3.24 (s, 3H),
10 3.10 (s, 6H), 2.70 (s, 3H), 2.68 (s, 3H), 1.54 (d, *J* = 9.7 Hz, 2H), 1.45 (s, 1H), 1.38–1.25 (m, 9H), 1.07–0.43 (m, 47H); ESI MS *m/z* 1365 [C₆₅H₁₁₀IN₁₁O₁₂ + H]⁺; HPLC >99% (AUC), *t*_R = 14.49 min.

Example 25 - Preparation of the Acetate of Cyclosporin Alkynyl Alcohol

15 [0089] A solution of acetyl cyclosporin vinyl iodide from Example 3 (75 mg, 0.055 mmol), CuI (11 mg, 0.055 mmol), and PdCl₂(PPh₃)₂ (39 mg, 0.055 mmol) in Et₃N (3 mL) was stirred at room temperature under N₂ atmosphere. Reaction mixture was then treated with 3-butyne-2-ol (39 mg, 0.55 mmol). Reaction was allowed to
20 stir for 6 h at room temperature. Solution was washed through micro filter and then concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford the acetate of cyclosporin alkynyl alcohol (15 mg, 21%) as an off-white solid:
¹H NMR (300 MHz, CDCl₃) δ 8.45 (d, *J* = 9.1 Hz, 1H), 7.99 (d, *J* = 6.1 Hz, 1H), 7.40 (d, *J* = 6.9 Hz, 2H), 5.72–5.13 (m, 12H), 5.01–4.39 (m, 10H), 4.26–4.07 (m, 2H), 3.45
25 (s, 3H), 3.37 (s, 3H), 3.26 (s, 3H), 3.21 (s, 3H), 3.10 (s, 3H), 2.68 (s, 3H), 2.66 (s, 3H), 2.04 (s, 6H), 1.50–1.43 (m, 5H), 1.31 (d, *J* = 7.8 Hz, 10H), 1.26 (s, 5H), 1.08–0.75 (m, 40H); ESI MS *m/z* 1299 [C₆₇H₁₁₅N₁₁O₁₄ + H]⁺.

Example 26 - Preparation of Cyclosporin Alkynyl Alcohol

30 [0090] A solution of acetate of cyclosporin alkynyl alcohol from Example 25 (15 mg, 0.011 mmol) in methanol (1 mL) was stirred at room temperature. Reaction mixture was treated with potassium carbonate (17 mg, 0.13 mmol) and allowed to keep stirring under N₂ atmosphere overnight. Mixture was diluted with ethyl acetate,

washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford cyclosporin alkynyl alcohol (5 mg, 36%) as a white solid: ¹H NMR (300 MHz, CDCl₃) δ 7.58 (d, *J* = 10.2 Hz, 1H), 7.41 (d, *J* = 8.0 Hz, 2H), 7.22 (d, *J* = 9.1 Hz, 1H), 6.06–5.94 (m, 2H), 5.66 (dd, *J* = 10.5, 3.2 Hz, 2H), 5.49 (s, 1H), 5.47–5.34 (m, 4H), 5.14–4.91 (s, 7H), 4.80–4.57 (m, 6H), 4.45 (t, *J* = 7.1 Hz, 2H), 4.04 (d, *J* = 7.0 Hz, 2H), 3.50 (s, 3H), 3.38 (s, 3H), 3.23 (s, 3H), 3.10 (s, 3H), 3.08 (s, 3H), 2.69 (s, 6H), 1.42 (d, *J* = 3.2 Hz, 6H), 1.38 (d, *J* = 7.2 Hz, 6H), 1.27–1.24 (m, 8H), 1.07–0.75 (m, 42H); ESI MS *m/z* 1257 [C₆₅H₁₁₃N₁₁O₁₃ + H]⁺; HPLC >99% (AUC), *t*_R = 13.44 min.

Example 27 – Preparation of Cyclosporin ene-yne-ene

[0091] The acetate of cyclosporin alkynyl alcohol from Example 25 (135 mg, 0.10 mmol) was dissolved in benzene (4 mL) and stirred under N₂ atmosphere. Solution was treated with Burgess reagent (98 mg, 0.40 mmol) and stirred at 60°C for 5 h. Mixture was diluted with ethyl ether, washed with water and brine, dried over sodium sulfate, and concentrated *in vacuo*.

[0092] The above crude product (68 mg, 0.053 mmol) was dissolved in methanol (3 mL) and stirred at room temperature. Reaction mixture was treated with potassium carbonate (60 mg, 0.43 mmol) and allowed to keep stirring under N₂ atmosphere overnight. Mixture was diluted with ethyl acetate, washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford cyclosporin ene-yne-ene (47 mg, 72%) as a white solid: ¹H NMR (300 MHz, CDCl₃) δ 7.92 (d, *J* = 9.8 Hz, 1H), 7.63 (d, *J* = 7.5 Hz, 1H), 7.42 (d, *J* = 8.2 Hz, 1H), 7.15 (d, *J* = 7.8 Hz, 1H), 6.16–6.10 (m, 2H), 5.92 (d, *J* = 2.0 Hz, 1H), 5.90 (d, *J* = 2.0 Hz, 1H), 5.86 (d, *J* = 2.0 Hz, 1H), 5.70 (dd, *J* = 10.8, 3.7 Hz, 2H), 5.63 (d, *J* = 2.1 Hz, 1H), 5.78–5.49 (m, 4H), 5.44 (d, *J* = 2.2 Hz, 1H), 5.41 (d, *J* = 2.2 Hz, 1H), 5.29 (dd, *J* = 11.4, 4.1 Hz, 2H), 5.11 (s, 2H), 5.09–4.91 (m, 4H), 4.80 (t, *J* = 7.3 Hz, 2H), 4.71–4.62 (m, 3H), 5.52 (d, *J* = 7.3 Hz, 2H), 3.51 (s, 3H), 3.39 (s, 3H), 3.25 (s, 3H), 3.10 (s, 6H), 2.70 (s, 3H), 2.68 (s, 3H), 1.35 (d, *J* = 7.2 Hz, 2H), 1.27–1.24 (m, 8H), 1.08–0.82 (m,

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45H), 0.73 (d, $J = 6.4$ Hz, 2H); ESI MS m/z 1239 [$C_{65}H_{111}N_{11}O_{12} + H$] $^{+}$; HPLC >99% (AUC), $t_R = 14.24$ min.

Example 28 - Preparation of Cyclosporin ene-yne-ene

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[0093] To a stirred solution of *trans*-acetyl cyclosporin vinyl iodide from Example 3 (130 mg, 0.095 mmol) in triethylamine (6 mL, degassed) were added 1-pentyn-3-ol (80 mg, 0.95 mmol), copper(I) iodide (18 mg, 0.095 mmol) and dichlorobis(triphenylphosphine)palladium(II) (67 mg, 0.095 mmol). After 18 h at room temperature under nitrogen, the mixture was filtered through a 0.2 μ m syringe filter and the solvent was removed under reduced pressure. The crude material was purified by flash column chromatography (SiO₂, 1:1 acetone/hexanes) to afford cyclosporin alkynyl alcohol (150 mg, ~80% pure by NMR) as a yellowish oil: ESI MS m/z 1313 [$C_{68}H_{117}N_{11}O_{14} + H$] $^{+}$.

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[0094] To a solution of the above cyclosporin alkynyl alcohol (150 mg, 0.11 mmol) in benzene (15 mL) under stirring and nitrogen was added Burgess reagent (135 mg, 0.57 mmol). The mixture was stirred at 60°C for 2 h. As MS monitoring still showed some starting material, some Burgess reagent (135 mg, 0.57 mmol) was added and the reaction was stirred 60°C for 2 h. After cooling down, the reaction was extracted with ethyl acetate (100 mL) and washed with water. The organic layer was separated, dried over anhydrous sodium sulfate, and concentrated under vacuum to afford the acetate of cyclosporin ene-yne-ene (38 mg), which was carried over without purification: ESI MS m/z 1295 [$C_{68}H_{115}N_{11}O_{13} + H$] $^{+}$.

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[0095] To a stirred solution of the above acetate of cyclosporin ene-yne-ene (38 mg, 0.029 mmol) in methanol (3 mL) was added potassium carbonate (30 mg, 0.21 mmol) at room temperature. After 18 h, ethyl acetate (100 mL) and water (50 mL) were added. The organic layer was separated, dried over anhydrous sodium sulfate, and concentrated under vacuum to afford crude product. The material was purified by semi-preparative HPLC to afford cyclosporin ene-yne-ene (3.1 mg, 8%) as a mixture of *cis* and *trans* isomers (yellow oil): 1H NMR (300 MHz, CDCl₃) δ 7.92 (d, $J = 9.5$ Hz, 1H), 7.63 (d, $J = 7.3$ Hz, 1H), 7.45 (d, $J = 8.0$ Hz, 1H), 7.14 (d, $J = 8.3$ Hz, 1H), 6.14–6.05 (m, 1H), 5.95–5.87 (m, 1H), 5.72–5.67 (m, 1H), 5.48–4.65 (m, 17H), 3.92–3.88 (m, 1H), 3.52 (s, 3H), 3.39 (s, 3H), 3.26 (s, 3H), 3.10 (s, 6H),

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2.70 (s, 3H), 2.68 (s, 3H), 2.70–0.70 (m, 67H); ESI MS m/z 1253 [$C_{66}H_{113}N_{11}O_{12} + H$]⁺; HPLC 13.6% and 86.3% (AUC), t_R = 15.49 and 16.67 min.

Example 29 - Preparation of Cyclosporin ene-yne-ene

- 5 [0096] To a stirred solution of *trans*-acetyl cyclosporin vinyl iodide from Example 3 (50 mg, 0.036 mmol) in triethylamine (4 mL, degassed) were added 2-methyl-1-buten-3-yne (24 mg, 0.36 mmol), copper(I) iodide (7 mg, 0.036 mmol) and dichlorobis(triphenylphosphine)palladium(II) (26 mg, 0.036 mmol). After 18 h at
- 10 room temperature under nitrogen, the mixture was filtered through a 0.2 μ m syringe filter and the solvent was removed under reduced pressure. The crude material was purified by semi-preparative HPLC to afford the acetate of cyclosporin ene-yne-ene (31 mg, 65%) as a yellowish oil: ESI MS m/z 1295 [$C_{68}H_{115}N_{11}O_{13} + H$]⁺.
- [0097] To a stirred solution of the above acetate of cyclosporin ene-yne-ene
- 15 (31 mg, 0.02 mmol) in methanol (3 mL) was added potassium carbonate (35 mg, 0.25 mmol) at room temperature. After 18 h, ethyl acetate (100 mL) and water (50 mL) were added. The organic layer was separated, dried over anhydrous sodium sulfate, and concentrated under vacuum to afford crude product. The material was purified by semi-preparative HPLC to afford cyclosporin ene-yne-ene (14 mg, 46%)
- 20 as a yellowish oil: ¹H NMR (500 MHz, CDCl₃) δ 7.93 (d, J = 9.5 Hz, 1H), 7.68–7.62 (m, 2H), 7.28 (d, J = 7.5 Hz, 1H), 6.17–6.11 (m, 1H), 6.08–6.03 (m, 1H), 5.65–5.58 (m, 2H), 5.56–5.44 (m, 2H), 5.42–4.50 (m, 8H), 3.92–3.90 (m, 1H), 3.78–3.73 (m, 1H), 3.50 (s, 3H), 3.38 (s, 3H), 3.24 (s, 3H), 3.11 (s, 3H), 3.09 (s, 3H), 2.72 (s, 3H), 2.70 (s, 3H), 1.89 (s, 3H), 2.45–0.71 (m, 69H); ESI MS m/z 1253 [$C_{66}H_{113}N_{11}O_{12} + H$]⁺; HPLC 96.4% (AUC), t_R = 16.55 min.
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Example 30 - Preparation of the Acetate of Cyclosporin Alkynyl Alcohol

- [0098] A mixture of acetyl cyclosporin vinyl iodide from Example 3 (70 mg, 0.051 mmol), CuI (5 mg, 0.026 mmol), and PdCl₂(PPh₃)₂ (18 mg, 0.026 mmol) in Et₃N (3 mL) was stirred at room temperature under N₂ atmosphere. Reaction mixture was then treated with 3-butyne-1-ol (37 mg, 0.51 mmol). Reaction was allowed to stir for 4 h at room temperature. Solution was filtered through micro-filter and then concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to
- 30

afford the acetate of cyclosporin alkynyl alcohol (52 mg, 78%) as an off-white solid: ^1H NMR (300 MHz, CDCl_3) δ 8.46 (d, $J = 9.5$ Hz, 1H), 8.04 (d, $J = 6.9$ Hz, 1H), 7.67 (d, $J = 9.2$ Hz, 1H), 7.46 (d, $J = 7.7$ Hz, 1H), 5.68 (dd, $J = 11.2, 4.6$ Hz, 2H), 5.58–4.78 (m, 18H), 4.63 (d, $J = 13.9$ Hz, 2H), 4.44 (t, $J = 6.9$ Hz, 2H), 3.98–3.77 (m, 4H), 3.45 (s, 3H), 3.24 (s, 3H), 3.23 (s, 3H), 3.19 (s, 3H), 3.08 (s, 3H), 2.67 (s, 3H), 2.66 (s, 3H), 2.02 (s, 2H), 1.30 (d, $J = 6.8$ Hz, 4H), 1.25 (t, $J = 2.2$ Hz, 4H), 1.05–0.75 (m, 52H); ESI MS m/z 1299 [$\text{C}_{67}\text{H}_{115}\text{N}_{11}\text{O}_{14} + \text{H}$] $^+$.

Example 31 - Preparation of Cyclosporin Alkynyl Alcohol

[0099] A solution of the acetate of cyclosporin alkynyl alcohol from Example 30 (52 mg, 0.04 mmol) in methanol (2 mL) was stirred at room temperature. Reaction mixture was treated with potassium carbonate (60 mg, 0.43 mmol) and was allowed to keep stirring under N_2 atmosphere overnight. Mixture was diluted with ethyl acetate, washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford cyclosporin alkynyl alcohol (22 mg, 44%) as a white solid: ^1H NMR (300 MHz, CDCl_3) δ 7.97 (d, $J = 9.7$ Hz, 1H), 7.62 (d, $J = 7.5$ Hz, 1H), 7.49 (d, $J = 8.4$ Hz, 1H), 7.20 (d, $J = 7.8$ Hz, 1H), 5.99–5.88 (m, 2H), 5.69 (dd, $J = 11.0, 4.0$ Hz, 1H), 5.44 (d, $J = 6.7$ Hz, 2H), 5.38 (s, 1H), 5.36 (d, $J = 3.6$ Hz, 1H), 5.32 (d, $J = 3.9$ Hz, 1H), 5.11–4.95 (m, 12H), 4.84 (t, $J = 7.1$ Hz, 2H), 4.72 (s, 1H), 4.68 (t, $J = 9.8$ Hz, 2H), 4.52 (t, $J = 7.2$ Hz, 1H), 3.89 (t, $J = 6.7$ Hz, 1H), 3.74 (t, $J = 5.7$ Hz, 1H), 3.49 (s, 3H), 3.40 (s, 3H), 3.26 (s, 3H), 3.11 (s, 3H), 3.10 (s, 3H), 2.69 (s, 3H), 2.68 (s, 3H), 1.35 (d, $J = 7.2$ Hz, 2H), 1.25 (d, $J = 2.9$ Hz, 4H), 1.07–0.70 (m, 54H); ESI MS m/z 1257 [$\text{C}_{65}\text{H}_{113}\text{N}_{11}\text{O}_{13} + \text{H}$] $^+$; HPLC >99% (AUC), $t_R = 17.67$ min.

Example 32 - Preparation of the Acetate of Cyclosporin Cyclopropyl Alkyne

[0100] A solution of cyclopropyl(trimethylsilyl)acetylene (76 mg, 0.55 mmol) in Et_3N (3 mL) was cooled to 0°C . Solution was treated with Bu_4NF (0.66 mL, 1 M in THF, 0.66 mmol) and allowed to stir at 0°C under N_2 atmosphere for 10 min. Reaction was allowed to warm to room temperature. After warming, the reaction mixture was treated with acetyl cyclosporin vinyl iodide from Example 3 (75 mg, 0.05 mmol), CuI (5 mg, 0.02 mmol), and $\text{PdCl}_2(\text{PPh}_3)_2$ (19 mg, 0.02 mmol). Reaction

was kept stirring for 3 h at room temperature. Solution was filtered through a micro filter and then concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford the acetate of cyclosporin cyclopropyl alkyne (58 mg, 82%) as an off-white solid: ^1H NMR (500 MHz, CDCl_3) δ 8.47 (d, $J = 4.5$ Hz, 1H), 8.01 (d, $J = 3.9$ Hz, 1H), 7.63 (d, $J = 5.4$ Hz, 1H), 7.57 (d, $J = 4.8$ Hz, 1H), 5.88–5.81 (m, 2H), 5.69 (dd, $J = 6.6, 2.4$ Hz, 2H), 5.58 (s, 1H), 5.54 (d, $J = 6.6$ Hz, 2H), 5.47 (s, 1H), 5.30–5.20 (m, 2H), 5.17 (d, $J = 3.6$ Hz, 2H), 5.15 (d, $J = 3.3$ Hz, 2H), 5.08–4.80 (m, 9H), 4.63 (s, 2H), 4.47 (t, $J = 4.2$ Hz, 2H), 4.13 (dd, $J = 8.7, 4.2$ Hz, 4H), 4.07 (t, $J = 3.9$ Hz, 2H), 3.42 (s, 3H), 3.25 (s, 3H), 3.22 (s, 3H), 3.19 (s, 3H), 3.09 (s, 3H), 2.69 (s, 3H), 2.67 (s, 3H), 1.32 (d, $J = 4.5$ Hz, 2H), 1.28 (dd, $J = 4.2, 1.6$ Hz, 4H), 1.10–0.71 (m, 51H); ESI MS m/z 1295 [$\text{C}_{68}\text{H}_{115}\text{N}_{11}\text{O}_{13} + \text{H}$] $^+$.

Example 33 - Preparation of Cyclosporin Cyclopropyl Alkyne

[0101] A solution of the acetate of cyclosporin cyclopropyl alkyne from Example 32 (58 mg, 0.044 mmol) in methanol (3 mL) was stirred at room temperature. Reaction mixture was treated with potassium carbonate (90 mg, 0.65 mmol) and allowed to keep stirring under N_2 atmosphere overnight. Mixture was diluted with ethyl acetate, washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated *in vacuo* to afford cyclosporin cyclopropyl alkyne (27 mg, 48%) as an off-white solid: ^1H NMR (300 MHz, CDCl_3) δ 7.90 (d, $J = 9.6$ Hz, 1H), 7.65 (d, $J = 7.3$ Hz, 1H), 7.50 (d, $J = 8.1$ Hz, 1H), 7.22 (d, $J = 7.8$ Hz, 1H), 6.03–5.92 (m, 2H), 5.70 (dd, $J = 10.7, 3.5$ Hz, 2H), 5.49 (d, $J = 5.7$ Hz, 2H), 5.38 (s, 1H), 5.33–4.99 (m, 12H), 4.93 (dd, $J = 9.5, 5.9$ Hz, 2H), 4.82 (d, $J = 5.7$ Hz, 2H), 4.76 (s, 2H), 4.71 (s, 2H), 4.65 (t, $J = 8.9$ Hz, 2H), 4.53 (t, $J = 7.2$ Hz, 2H), 3.49 (s, 3H), 3.37 (s, 3H), 3.24 (s, 3H), 3.10 (s, 6H), 2.71 (s, 3H), 2.69 (s, 3H), 1.35 (d, $J = 7.2$ Hz, 2H), 1.26 (d, $J = 6.6$ Hz, 6H), 1.07–0.48 (m, 49H); ESI MS m/z 1253 [$\text{C}_{66}\text{H}_{113}\text{N}_{11}\text{O}_{12} + \text{H}$] $^+$; HPLC >99% (AUC), $t_R = 20.26$ min.

Example 34 – Preparation of the Acetate of Cyclosporin Alkyne

[0102] A mixture of acetyl cyclosporin vinyl iodide from Example 3 (100 mg, 0.073 mmol), CuI (7 mg, 0.037 mmol), and $\text{PdCl}_2(\text{PPh}_3)_2$ (26 mg, 0.037 mmol) in Et_3N (2 mL) was stirred at room temperature under N_2 atmosphere. Reaction mixture

was then treated with cyclohexylacetylene (80 mg, 0.737 mmol). Reaction was allowed to stir for 6 h at room temperature. Solution was filtered through a short pad of silica gel and then concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford the acetate of cyclosporin alkyne (75 mg, 76%) as a light brown solid: ^1H NMR (300 MHz, CDCl_3) δ 8.43 (d, $J = 9.4$ Hz, 1H), 8.07 (d, $J = 6.7$ Hz, 1H), 7.85 (d, $J = 11.7$ Hz, 1H), 7.66 (d, $J = 11.4$ Hz, 1H), 6.00–5.90 (m, 1H), 5.80–5.60 (m, 2H), 5.55–5.45 (m, 2H), 5.40–5.20 (m, 3H), 5.10–4.80 (m, 3H), 4.70–4.45 (m, 2H), 3.42 (s, 3H), 3.24 (s, 3H), 3.21 (m, 3H), 3.19 (s, 3H), 3.11 (s, 3H), 2.70 (s, 3H), 2.69 (s, 3H), 2.20–0.80 (m, 83H); ESI MS m/z 1337 [$\text{C}_{71}\text{H}_{121}\text{N}_{11}\text{O}_{13} + \text{H}$] $^+$.

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Example 35 - Preparation of Cyclosporin Alkyne

[0103] A solution of the acetate of cyclosporin alkyne from Example 34 (76 mg, 0.057 mmol) in methanol (2 mL) was stirred at room temperature. Reaction mixture was treated with potassium carbonate (40 mg, 0.29 mmol) and allowed to keep stirring under N_2 atmosphere overnight. Mixture was diluted with ethyl acetate, washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford cyclosporin alkyne (29 mg, 39%) as a white solid: ^1H NMR (300 MHz, CDCl_3) δ 7.94 (d, $J = 9.8$ Hz, 1H), 7.65–7.55 (m, 2H), 7.26 (overlapped with CHCl_3 , 1H), 6.00–5.85 (m, 1H), 5.71 (dd, $J = 11.0, 4.0$ Hz, 1H), 5.53 (d, $J = 5.5$ Hz, 1H), 5.40 (d, $J = 15.6$ Hz, 1H), 5.30 (dd, $J = 11.3, 3.5$ Hz, 1H), 5.13 (d, $J = 10.8$ Hz, 1H), 5.10–5.00 (m, 3H), 4.89 (dd, $J = 9.2, 6.2$ Hz, 1H), 4.83 (t, $J = 7.1$ Hz, 1H), 4.73 (d, $J = 14.1$ Hz, 1H), 4.63 (t, $J = 9.7$ Hz, 1H), 4.56 (t, $J = 7.3$ Hz, 1H), 3.73 (t, $J = 7.3$ Hz, 1H), 3.50 (s, 3H), 3.38 (s, 3H), 3.25 (s, 3H), 3.11 (s, 3H), 3.09 (s, 3H), 2.71 (s, 3H), 2.70 (s, 3H), 2.60–2.55 (m, 1H), 2.45–2.35 (m, 2H), 2.20–0.75 (m, 74H), 0.68 (d, $J = 5.8$ Hz, 2H); ESI MS m/z 1295 [$\text{C}_{69}\text{H}_{119}\text{N}_{11}\text{O}_{12} + \text{H}$] $^+$; HPLC 97.9 % (AUC), $t_R = 23.24$ min.

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Example 36 - Preparation of Cyclosporin Alkyne

[0104] To a stirred solution of *trans*-acetyl cyclosporin vinyl iodide from Example 3 (50 mg, 0.036 mmol) in triethylamine (3 mL, degassed) were added phenylacetylene (37 mg, 0.36 mmol), copper(I) iodide (7.0 mg, 0.036 mmol) and

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dichlorobis(triphenylphosphine)palladium(II) (26 mg, 0.036 mmol). After 18 h at room temperature under nitrogen, the mixture was filtered through a 0.2 μ m syringe filter and the solvent was removed under reduced pressure. The crude material was purified by semi-preparative HPLC to afford the acetate of cyclosporin alkyne (34 mg, 69%) as a yellowish oil: ESI MS m/z 1331 [$C_{71}H_{115}N_{11}O_{13} + H$] $^{+}$.

- 5 [0105] To a stirred solution of the above acetate of cyclosporin alkyne (34 mg, 0.02 mmol) in methanol (3 mL) was added potassium carbonate (30 mg, 0.2 mmol) at room temperature. After 18 h, ethyl acetate (100 mL) and water (50 mL) were added. The organic layer was separated, dried over anhydrous sodium sulfate, and
- 10 concentrated under vacuum to afford crude product. The material was purified by semi-preparative HPLC to afford cyclosporin alkyne (15 mg, 45%) as a yellowish oil: 1H NMR (500 MHz, $CDCl_3$) δ 7.92 (m, 1H), 7.61 (m, 1H), 7.49 (m, 1H), 7.41 (m, 3H), 7.21 (m, 1H), 6.20 (m, 1H), 5.65–4.45 (m, 20H), 3.98 (m, 1H), 3.73 (m, 1H), 3.52 (s, 3H), 3.39 (s, 3H), 3.25 (s, 3H), 3.10 (s, 6H), 2.71 (s, 3H), 2.69 (s, 3H),
- 15 2.45–0.70 (m, 62H). ESI MS m/z 1289 [$C_{69}H_{113}N_{11}O_{12} + H$] $^{+}$; HPLC >98% (AUC); t_R = 16.67 min.

Example 37 - Preparation of the Acetate of Cyclosporin Alkyne

- 20 [0106] A mixture of acetyl cyclosporin vinyl iodide from Example 3 (80 mg, 0.059 mmol), CuI (6 mg, 0.030 mmol), and $PdCl_2(PPh_3)_2$ (21 mg, 0.030 mmol) in Et_3N (2 mL) was stirred at room temperature under N_2 atmosphere. To the above reaction mixture was then treated with 1-chloro-4-ethynylbenzene (80 mg, 0.59 mmol). The resulting reaction mixture was stirred overnight at room
- 25 temperature. Reaction was filtered through a short pad of silica gel and filtrate was concentrated under reduced pressure. The crude product was purified by semi-preparative HPLC to afford the acetate of cyclosporin alkyne (54 mg, 67%) as a light brown solid: 1H NMR (300 MHz, $CDCl_3$) δ 8.47 (d, J = 9.6 Hz, 1H), 8.06 (d, J = 6.8 Hz, 1H), 7.85–7.65 (m, 2H), 7.61 (d, J = 7.6 Hz, 1H), 7.45–7.35 (m, 1H), 7.33–
- 30 7.27 (m, 2H), 6.05–5.95 (m, 1H), 5.69 (dd, J = 11.0, 3.9 Hz, 1H), 5.60–5.47 (m, 2H), 5.45–5.35 (m, 1H), 5.29 (dd, J = 8.7, 4.0 Hz, 1H), 5.20–5.10 (m, 1H), 5.05–4.70 (m, 4H), 4.63 (d, J = 14.1 Hz, 1H), 4.46 (t, J = 7.0 Hz, 1H), 3.44 (s, 3H), 3.27 (s, 3H),

3.26 (s, 3H), 3.20 (s, 3H), 3.09 (s, 3H), 2.69 (s, 3H), 2.68 (s, 3H), 2.50–2.40 (m, 1H), 2.30–0.70 (m, 71H); ESI MS m/z 1365 [$C_{71}H_{114}ClN_{11}O_{13} + H$]⁺.

Example 38 - Preparation of Cyclosporin Alkyne

5 **[0107]** A solution of the acetate of cyclosporin alkyne from Example 37 (54 mg, 0.047 mmol) in methanol (1.5 mL) was stirred at room temperature. Reaction mixture was treated with potassium carbonate (30 mg, 0.21 mmol) and allowed to keep stirring under N₂ atmosphere overnight. Mixture was diluted with ethyl acetate,
10 washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated under reduced pressure. The crude product was purified by semi-preparative HPLC to afford cyclosporin alkyne (32 mg, 62%) as an off-white solid: ¹H NMR (500 MHz, CDCl₃) δ 7.96 (d, J = 9.7 Hz, 1H), 7.65 (d, J = 7.4 Hz, 1H), 7.56 (d, J = 8.2 Hz, 1H), 7.40–7.33 (m, 2H), 7.30–7.20 (m, 3H), 6.22–6.15 (m
15 1H), 5.70 (dd, J = 11.0, 4.2 Hz, 1H), 5.62 (d, J = 15.8 Hz, 1H), 5.52 (d, J = 6.2 Hz, 1H), 5.32 (dd, J = 11.4, 3.6 Hz, 1H), 5.12–5.00 (m, 2H), 4.96 (dd, J = 9.8, 5.9 Hz, 2H), 4.82 (t, J = 7.3 Hz, 1H), 4.72 (d, J = 14.0 Hz, 1H), 4.67 (t, J = 8.9 Hz, 1H), 4.54 (t, J = 7.3 Hz, 1H), 3.82 (t, J = 6.7 Hz, 1H), 3.52 (s, 3H), 3.40 (s, 3H), 3.26 (s, 3H), 3.11 (s, 3H), 3.10 (s, 3H), 2.71 (s, 3H), 2.69 (s, 3H), 2.65–2.55 (m, 1H), 2.45–2.35
20 (m, 1H), 2.20–0.85 (m, 65H), 0.75 (d, J = 6.9 Hz, 2H); ESI MS m/z 1323 [$C_{69}H_{112}ClN_{11}O_{12} + H$]⁺; HPLC 97.3% (AUC), t_R = 22.54 min.

Example 39 - Preparation of the Acetate of Cyclosporin Alkyne

25 **[0108]** A mixture of acetyl cyclosporin vinyl iodide from Example 3 (75 mg, 0.055 mmol), CuI (6 mg, 0.030 mmol), and PdCl₂(PPh₃)₂ (21 mg, 0.030 mmol) in Et₃N (2 mL) was stirred at room temperature under N₂ atmosphere. Reaction mixture was then treated with 3-ethynylthiophene (65 mg, 0.60 mmol). Reaction was allowed to stir for 6 h at room temperature. Solution was filtered through a short pad of silica
30 gel and then concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford the acetate of cyclosporin alkyne (31 mg, 42%) as a light brown solid: ¹H NMR (300 MHz, CDCl₃) δ 8.47 (d, J = 9.5 Hz, 1H), 8.06 (d, J = 6.7 Hz, 1H), 7.73 (d, J = 9.2 Hz, 1H), 7.63 (d, J = 7.6 Hz, 1H), 7.49 (d, J = 2.9 Hz, 1H), 7.25–7.20 (m, 1H), 7.15 (d, J = 5.0 Hz, 1H), 6.10–5.90 (m, 1H), 5.70 (dd, J =

11.2, 3.5 Hz, 1H), 5.60–4.50 (m, 9H), 4.63 (d, $J = 14.2$ Hz, 1H), 4.51 (t, $J = 7.2$ Hz, 1H), 3.43 (s, 3H), 3.26 (s, 6H), 3.20 (s, 3H), 3.09 (s, 3H), 2.72 (s, 3H), 2.68 (s, 3H), 2.50–2.40 (m, 1H), 2.30–0.70 (m, 71H); ESI MS m/z 1337 [$C_{69}H_{113}N_{11}O_{13}S + H$]⁺.

5 **Example 40 - Preparation of Cyclosporin Alkyne**

[0109] A solution of the acetate of cyclosporin alkyne from Example 39 (31 mg, 0.023 mmol) in methanol (2 mL) was stirred at room temperature. Reaction mixture was treated with potassium carbonate (15 mg, 0.108 mmol) and allowed to
10 keep stirring under N₂ atmosphere overnight. Mixture was diluted with ethyl acetate, washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford cyclosporin alkyne (15 mg, 50%) as a white solid: ¹H NMR (500 MHz, CDCl₃) δ 7.94 (d, $J = 9.8$ Hz, 1H), 7.66 (d, $J = 7.3$ Hz, 1H), 7.60 (d, $J = 8.2$ Hz, 1H), 7.50 (dd, $J = 3.0, 1.0$ Hz, 1H), 7.30 (d, $J = 7.8$ Hz, 1H), 7.24 (dd, $J = 5.0, 3.1$ Hz, 1H), 7.11 (dd, $J = 5.0, 1.1$ Hz, 1H), 6.30–6.20 (m, 1H), 5.70 (dd, $J = 11.0, 4.2$ Hz, 1H), 5.61 (d, $J = 15.8$ Hz, 1H), 5.51 (d, $J = 6.2$ Hz, 1H), 5.34 (dd, $J = 11.3, 3.5$ Hz, 1H), 5.15–5.00 (m, 3H), 4.95 (dd, $J = 9.5, 5.9$ Hz, 1H), 4.83 (t, $J = 7.2$ Hz, 1H), 4.73 (d, $J = 14.1$ Hz, 1H), 4.67 (t, $J = 9.5$ Hz, 1H), 4.54 (t, $J = 7.3$ Hz, 1H), 3.83
15 (t, $J = 6.9$ Hz, 1H), 3.51 (s, 3H), 3.39 (s, 3H), 3.24 (s, 3H), 3.11 (s, 3H), 3.10 (s, 3H), 2.72 (s, 3H), 2.70 (s, 3H), 2.65–2.55 (m, 1H), 2.50–2.40 (m, 1H), 2.20–0.84 (m, 67H); ESI MS m/z 1295 [$C_{67}H_{111}N_{11}O_{12}S + H$]⁺; HPLC 97.9% (AUC), $t_R = 20.46$ min.

25 **Example 41 - Preparation of the Acetate of Cyclosporin Alkyne**

[0110] A mixture of acetyl cyclosporin vinyl iodide from Example 3 (80 mg, 0.059 mmol), CuI (6 mg, 0.030 mmol), and PdCl₂(PPh₃)₂ (21 mg, 0.030 mmol) in Et₃N (2 mL) was stirred at room temperature under N₂ atmosphere. Reaction mixture was then treated with 2-ethynylpyridine (61 mg, 0.59 mmol). Reaction was allowed
30 to keep stirring under nitrogen atmosphere at room temperature overnight. The reaction mixture was filtered through a short pad of silica gel and then concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford the acetate of cyclosporin alkyne (50 mg, 64%) as an off-white solid: ¹H NMR (300 MHz, CDCl₃) δ 8.56 (d, $J = 4.8$ Hz, 1H), 8.52 (d, $J = 9.7$ Hz, 1H), 8.03 (d, $J =$

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6.9 Hz, 1H), 7.70–7.60 (m, 2H), 7.50–7.40 (m, 2H), 7.30–7.20 (m, 1H), 6.20–6.10 (m, 1H), 5.68 (dd, $J = 9.0, 4.0$ Hz, 1H), 5.62–5.50 (m, 3H), 5.40–5.25 (m, 2H), 5.20–5.10 (m, 1H), 5.05–4.95 (m, 2H), 4.85–4.75 (m, 1H), 4.64 (d, $J = 13.8$ Hz, 1H), 4.42 (t, $J = 7.1$ Hz, 1H), 3.45 (s, 3H), 3.30 (s, 3H), 3.28 (s, 3H), 3.21 (s, 3H), 3.07 (s, 3H), 2.68 (s, 3H), 2.66 (s, 3H), 2.50–2.40 (m, 2H), 2.30–0.70 (m, 71H); ESI MS m/z 1332 [$C_{70}H_{114}N_{12}O_{13} + H$] $^{+}$.

Example 42 - Preparation of Cyclosporin Alkyne

10 [0111] A solution of the acetate of cyclosporin alkyne from Example 41 (50 mg, 0.0375 mmol) in methanol (2 mL) was stirred at room temperature. Reaction mixture was treated with potassium carbonate (35 mg, 0.25 mmol) and allowed to keep stirring under N_2 atmosphere overnight. Mixture was diluted with ethyl acetate, washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated under reduced pressure. The crude product was purified by semi-preparative HPL to afford cyclosporin alkyne (3 mg, 6%) as a white solid; 1H NMR (500 MHz, $CDCl_3$) δ 8.79 (d, $J = 4.7$ Hz, 1H), 8.00–7.90 (m, 2H), 7.71 (d, $J = 7.9$ Hz, 1H), 7.65 (d, $J = 7.6$ Hz, 1H), 7.50–7.30 (m, 2H), 7.16 (d, $J = 7.9$ Hz, 1H), 6.50–6.40 (m, 1H), 5.75–5.65 (m, 2H), 5.53 (d, $J = 6.2$ Hz, 1H), 5.30–5.25 (m, 1H), 5.15–5.00 (m, 3H), 4.96 (dd, $J = 9.9, 3.5$ Hz, 1H), 4.83 (t, $J = 7.2$ Hz, 1H), 4.72 (d, $J = 13.9$ Hz, 1H), 4.68 (t, $J = 9.2$ Hz, 1H), 4.51 (t, $J = 7.0$ Hz, 1H), 3.84 (t, $J = 6.8$ Hz, 1H), 3.52 (s, 3H), 3.40 (s, 3H), 3.26 (s, 3H), 3.11 (s, 3H), 3.10 (s, 3H) 2.71 (s, 3H), 2.69 (s, 3H), 2.65–2.60 (m, 1H), 2.45–2.40 (m, 1H), 2.20–0.70 (m, 67H); ESI MS m/z 1290 [$C_{68}H_{112}N_{12}O_{12} + H$] $^{+}$; HPLC >99% (AUC), $t_R = 18.94$ min.

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Example 43 - Preparation of the Acetate of Cyclosporin Alkyne

[0112] A mixture of acetyl cyclosporin vinyl iodide from Example 3 (100 mg, 0.073 mmol), CuI (7 mg, 0.037 mmol), and $PdCl_2(PPh_3)_2$ (26 mg, 0.037 mmol) in Et_3N (2 mL) was stirred at room temperature under N_2 atmosphere. Reaction mixture was then treated with 1-pentyne (50 mg, 0.737 mmol). Reaction was allowed to stir for 6 h at room temperature. Solution was filtered through a short pad of silica gel then concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford the acetate of cyclosporin alkyne (51 mg, 54%) as an off-white solid:

¹H NMR (300 MHz, CDCl₃) δ 8.43 (d, *J* = 9.4 Hz, 1H), 8.06 (d, *J* = 6.7 Hz, 1H), 7.83 (d, *J* = 9.3 Hz, 1H), 7.64 (d, *J* = 7.6 Hz, 1H), 6.00–4.40 (m, 13H), 3.43 (s, 3H), 3.24 (s, 3H), 3.23 (s, 3H), 3.19 (s, 3H), 3.11 (s, 3H), 2.70 (3H), 2.69 (s, 3H), 2.45–0.75 (m, 79H); ESI MS *m/z* 1297 [C₆₈H₁₁₇N₁₁O₁₃ + H]⁺.

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Example 44 - Preparation of Cyclosporin Alkyne

[0113] A solution of the acetate of cyclosporin alkyne from Example 43 (51 mg, 0.039 mmol) in methanol (2 mL) was stirred at room temperature. Reaction mixture was treated with potassium carbonate (26 mg, 0.18 mmol) and allowed to keep stirring under N₂ atmosphere overnight. Mixture was diluted with ethyl acetate, washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford cyclosporin alkyne (35 mg, 71%) as a white solid: ¹H NMR (300 MHz, CDCl₃) δ 7.91 (d, *J* = 9.8 Hz, 1H), 7.63 (d, *J* = 7.5 Hz, 1H), 7.48 (d, *J* = 8.2 Hz, 1H), 7.18 (d, *J* = 7.8 Hz, 1H), 6.10–5.95 (m, 1H), 5.71 (dd, *J* = 13.0, 4.1 Hz, 1H), 5.51 (d, *J* = 5.7 Hz, 1H), 5.40 (d, *J* = 15.8 Hz, 1H), 5.30 (dd, *J* = 11.5, 4.0 Hz, 1H), 5.12 (d, *J* = 10.9 Hz, 1H), 5.07–4.95 (m, 2H), 4.92 (dd, *J* = 9.6, 6.0 Hz, 1H), 4.82 (t, *J* = 7.3 Hz, 1H), 4.73 (d, *J* = 14.1 Hz, 1H), 4.64 (t, *J* = 8.3 Hz, 1H), 4.54 (t, *J* = 7.3 Hz, 1H), 3.76 (t, *J* = 6.8 Hz, 1H), 3.51 (s, 3H), 3.38 (s, 3H), 3.25 (s, 3H), 3.09 (s, 6H), 2.71 (s, 3H), 2.69 (s, 3H), 2.65–2.55 (m, 1H), 2.45–2.35 (m, 1H), 2.30–2.20 (m, 2H), 2.15–1.95 (m, 4H), 1.85–0.80 (m, 66H), 0.69 (d, *J* = 6.0 Hz, 2H); ESI MS *m/z* 1255 [C₆₆H₁₁₅N₁₁O₁₂ + H]⁺; HPLC >99% (AUC), *t_R* = 20.98 min.

Example 45 - Preparation of the Acetate of Cyclosporin Alkyne

[0114] A mixture of acetyl cyclosporin vinyl iodide from Example 3 (100 mg, 0.073 mmol), CuI (7 mg, 0.037 mmol), and PdCl₂(PPh₃)₂ (26 mg, 0.037 mmol) in Et₃N (2 mL) was stirred at room temperature under N₂ atmosphere. Reaction mixture was then treated with 1-hexyne (80 mg, 0.737 mmol). Reaction was allowed to stir for 4 h at room temperature. Solution was filtered through a short pad of silica gel and then concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford the acetate of cyclosporin alkyne (61 mg, 63%) as a light brown solid: ¹H NMR (300 MHz, CDCl₃) δ 8.45 (d, *J* = 9.6 Hz, 1H), 8.05 (d, *J* = 6.8 Hz, 1H), 7.66

(d, $J = 9.2$ Hz, 1H), 7.57 (d, $J = 7.7$ Hz, 1H), 5.90–6.00 (m, 1H), 5.69 (dd, $J = 10.9$, 3.8 Hz, 1H), 5.60–5.10 (m, 5H), 5.05–4.75 (m, 4H), 4.68 (d, $J = 13.3$ Hz, 1H), 3.44 (s, 3H), 3.25 (s, 3H), 3.23 (s, 3H), 3.19 (s, 3H), 3.09 (s, 3H), 2.69 (s, 3H), 2.67 (s, 3H), 2.45–0.75 (m, 82H); ESI MS m/z 1311 [$C_{69}H_{119}N_{11}O_{13} + H$] $^{+}$.

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Example 46 - Preparation of Cyclosporin Alkyne

[0115] To a solution of the acetate of cyclosporin alkyne from Example 45 (61 mg, 0.047 mmol) in methanol (2 mL) was added potassium carbonate (30 mg, 0.22 mmol) at room temperature and allowed to keep stirring under N_2 atmosphere overnight. Mixture was diluted with ethyl acetate, washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford cyclosporin alkyne (28 mg, 49%) as a white solid: 1H NMR (300 MHz, $CDCl_3$) δ 7.91 (d, $J =$ 9.7 Hz, 1H), 7.62 (d, $J = 7.8$ Hz, 1H), 7.48 (d, $J = 8.2$ Hz, 1H), 7.18 (d, $J = 7.9$ Hz, 1H), 6.10–5.90 (m, 1H), 5.71 (dd, $J = 11.0$, 3.9 Hz, 1H), 5.51 (d, $J = 5.7$ Hz, 1H), 5.40 (d, $J = 15.7$ Hz, 1H), 5.30 (dd, $J = 11.5$, 3.9 Hz, 1H), 5.12 (d, $J = 10.9$ Hz, 1H), 5.07–4.95 (m, 2H), 4.92 (dd, $J = 9.6$, 6.0 Hz, 1H), 4.80 (t, $J = 7.4$ Hz, 1H), 4.74 (d, $J =$ 14.1 Hz, 1H), 4.64 (t, $J = 8.3$ Hz, 1H), 4.54 (t, $J = 7.3$ Hz, 1H), 3.76 (t, $J = 6.8$ Hz, 1H), 3.51 (s, 3H), 3.38 (s, 3H), 3.26 (s, 3H), 3.10 (s, 6H), 2.71 (s, 3H), 2.69 (s, 3H), 2.60–2.50 (m, 1H), 2.45–2.35 (m, 1H), 2.33–2.22 (m, 2H), 2.20–2.00 (m, 4H), 1.85–0.80 (m, 68H), 0.69 (d, $J = 5.8$ Hz, 2H); ESI MS m/z 1269 [$C_{67}H_{117}N_{11}O_{12} + H$] $^{+}$; HPLC 98.4% (AUC), $t_R = 21.91$ min.

Example 47 - Preparation of the Acetate of Cyclosporin Alkyne

[0116] A mixture of acetylcyclosporin vinyl iodide from Example 3 (80 mg, 0.059 mmol), CuI (6 mg, 0.030 mmol), and $PdCl_2(PPh_3)_2$ (21 mg, 0.030 mmol) in Et_3N (2 mL) was stirred at room temperature under N_2 atmosphere. Reaction mixture was then treated with 5-chloro-1-pentyne (60 mg, 0.59 mmol). Reaction was allowed to stir for 6 h at room temperature. Solution was filtered through a short pad of silica gel and then concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford the acetate of cyclosporin alkyne (76 mg, 97%) as a light brown solid: 1H NMR (300 MHz, $CDCl_3$) δ 8.44 (d, $J = 9.5$ Hz, 1H), 8.05 (d, $J =$

6.8 Hz, 1H), 7.74 (d, $J = 9.3$ Hz, 1H), 7.61 (d, $J = 7.7$ Hz, 1H), 5.95–5.80 (m, 1H), 5.69 (dd, $J = 11.1, 3.8$ Hz, 1H), 5.60–5.45 (m, 3H), 5.35–5.10 (m, 2H), 5.05–4.70 (m, 4H), 4.64 (d, $J = 14.0$ Hz, 1H), 4.46 (t, $J = 7.1$ Hz, 1H), 3.64 (t, $J = 6.4$ Hz, 1H), 3.44 (s, 3H), 3.25 (s, 3H), 3.23 (s, 3H), 3.19 (s, 3H), 3.10 (s, 3H), 2.69 (s, 3H), 2.68 (s, 3H), 2.70–2.60 (m, 2H), 2.25–1.50 (m, 17 h), 1.45–0.60 (m, 58H); m/z 1331 [$C_{68}H_{116}ClN_{11}O_{13} + H$] $^{+}$.

Example 48 - Preparation of Cyclosporin Alkyne

10 [0117] A solution of the acetate of cyclosporin alkyne from Example 47 (76 mg, 0.047 mmol) in methanol (2 mL) was stirred at room temperature. Reaction mixture was treated with potassium carbonate (35 mg, 0.25 mmol) and allowed to keep stirring under N_2 atmosphere overnight. Mixture was diluted with ethyl acetate, washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford cyclosporin alkyne (29 mg, 49%) as an off-white solid:

15 1H NMR (500 MHz, $CDCl_3$) δ 7.92 (d, $J = 9.8$ Hz, 1H), 7.61 (d, $J = 7.5$ Hz, 1H), 7.49 (d, $J = 8.2$ Hz, 1H), 7.19 (d, $J = 7.9$ Hz, 1H), 6.10–5.95 (m, 1H), 5.70 (dd, $J = 11.0, 4.3$ Hz, 1H), 5.51 (d, $J = 5.9$ Hz, 1H), 5.39 (d, $J = 16.1$ Hz, 1H), 5.29 (dd, $J = 11.4, 3.8$ Hz, 1H), 5.11 (d, $J = 10.9$ Hz, 1H), 5.07–5.00 (m, 2H), 4.92 (dd, $J = 9.6, 6.0$ Hz, 1H), 4.81 (t, $J = 7.2$ Hz, 1H), 4.73 (d, $J = 14.0$ Hz, 1H), 4.64 (dd, $J = 9.6, 8.6$ Hz, 1H), 4.54 (t, $J = 7.3$ Hz, 1H), 3.77 (t, $J = 7.1$ Hz, 1H), 3.64 (t, $J = 3.4$ Hz, 3H), 3.51 (s, 3H), 3.39 (s, 3H), 3.25 (s, 3H), 3.11 (s, 3H), 3.10 (s, 3H), 2.71 (s, 3H), 2.69 (s, 3H), 2.60–2.55 (m, 1H), 2.50–2.40 (m, 3H), 2.20–0.80 (m, 66 h), 0.70 (d, $J = 6.3$ Hz, 2H); m/z

20 1289 [$C_{66}H_{114}ClN_{11}O_{12} + H$] $^{+}$; HPLC 98.1 % (AUC), $t_R = 20.39$ min.

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Example 49 – Preparation of the Acetate of Cyclosporin Alkynyl Ether

[0118] A solution of the acetate of cyclosporin alkynyl alcohol from Example 20 (77 mg, 0.06 mmol) in methylene chloride (2 mL) was stirred at room temperature under N_2 atmosphere. Solution was treated with allyl chloride (46 mg, 0.60 mmol), $PhCH_2NEt_3Cl$ (14 mg, 0.06 mmol) and 40% KOH in water (2 mL). The reaction was stirred for 6 h. Mixture was diluted with ether and washed with water and brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product

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was purified by semi-preparative HPLC to afford the acetate of cyclosporin alkynyl ether (18 mg, 23%) as an off-white solid: ^1H NMR (300 MHz, CDCl_3) δ 8.43 (d, $J = 9.6$ Hz, 1H), 8.01 (d, $J = 6.6$ Hz, 1H), 7.61 (d, $J = 9.1$ Hz, 1H), 7.55 (d, $J = 7.7$ Hz, 1H), 5.98–5.86 (m, 4H), 5.52 (d, $J = 9.8$ Hz, 2H), 5.38–4.62 (m, 17H), 4.45 (d, $J = 6.8$ Hz, 1H), 4.41 (d, $J = 1.2$ Hz, 2H), 4.26 (s, 2H), 4.13 (d, $J = 2.1$ Hz, 2H), 4.06 (d, $J = 5.5$ Hz, 2H), 3.43 (s, 3H), 3.25 (s, 3H), 3.23 (s, 3H), 3.19 (s, 3H), 3.09 (s, 3H), 2.68 (s, 3H), 2.66 (s, 3H), 1.54 (d, $J = 10.2$ Hz, 2H), 1.46 (s, 2H), 1.37 (s, 2H), 1.38–1.19 (m, 4H), 1.06–0.84 (m, 48H), 0.80 (d, $J = 6.5$ Hz, 2H); ESI MS m/z 1325 $[\text{C}_{69}\text{H}_{117}\text{N}_{11}\text{O}_{14} + \text{H}]^+$.

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Example 50 - Preparation of Cyclosporin Alkynyl Ether

[0119] A solution of the acetate of cyclosporin alkynyl ether from Example 49 (14.5 mg, 0.01 mmol) in methanol (2 mL) was stirred at room temperature. Reaction mixture was treated with potassium carbonate (25 mg, 0.18 mmol) and was allowed to keep stirring under N_2 atmosphere overnight. Mixture was diluted with ethyl acetate, washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford cyclosporin alkynyl ether (6 mg, 42%) as a white solid:

20 ^1H NMR (300 MHz, CDCl_3) δ 7.94 (d, $J = 9.6$ Hz, 1H), 7.62 (d, $J = 7.6$ Hz, 1H), 7.44 (d, $J = 8.4$ Hz, 1H), 7.14 (d, $J = 7.9$ Hz, 1H), 6.12–5.68 (m, 2H), 5.51 (d, $J = 5.7$ Hz, 2H), 5.47 (s, 1H), 5.42 (s, 1H), 5.35–5.28 (m, 1H), 5.22 (d, $J = 1.3$ Hz, 2H), 5.19 (d, $J = 1.2$ Hz, 2H), 5.12 (s, 2H), 5.08 (s, 2H), 5.07–4.81 (m, 2H), 4.75 (s, 1H), 4.70 (s, 1H), 4.26 (d, $J = 1.7$ Hz, 3H), 4.13 (s, 2H), 4.12–4.04 (m, 2H), 3.51 (s, 3H), 3.39 (s, 3H), 3.26 (s, 3H), 3.11 (s, 3H), 3.10 (s, 3H), 2.70 (s, 3H), 2.68 (s, 3H), 2.13 (s, 2H), 2.04 (s, 3H), 1.53 (d, $J = 6.9$ Hz, 1H), 1.45 (s, 2H), 1.37 (s, 2H), 1.34 (d, $J = 10.8$ Hz, 2H), 1.25 (s, 2H), 1.09–0.81 (m, 48H), 0.71 (d, $J = 6.1$ Hz, 2H); ESI MS m/z 1283 $[\text{C}_{67}\text{H}_{115}\text{N}_{11}\text{O}_{13} + \text{H}]^+$; HPLC 95.1% (AUC), $t_R = 19.71$ min.

Example 51 - Preparation of the Acetate of Cyclosporin Alkynyl Ether

[0120] A mixture of acetyl cyclosporin vinyl iodide from Example 3 (70 mg, 0.051 mmol), CuI (5 mg, 0.026 mmol) and $\text{PdCl}_2(\text{PPh}_3)_2$ (18 mg, 0.026 mmol) in Et_3N (3 mL) was stirred at room temperature under N_2 atmosphere. Reaction mixture

was then treated with methyl propargyl ether (36 mg, 0.51 mmol). Reaction was allowed to stir for 4 h at room temperature. Solution was filtered through micro-filter and then concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford the acetate of cyclosporin alkynyl ether (61 mg, 91%) as an off-white solid: ^1H NMR (300 MHz, CDCl_3) δ 8.39 (d, $J = 9.4$ Hz, 1H), 8.02 (d, $J = 6.5$ Hz, 1H), 7.70 (d, $J = 9.1$ Hz, 1H), 7.69 (d, $J = 8.6$ Hz, 1H), 6.04–5.91 (m, 4H), 5.52 (d, $J = 11.1$ Hz, 2H), 5.39–5.14 (m, 8H), 5.06–4.61 (m, 12H), 4.45 (t, $J = 7.0$ Hz, 2H), 4.21 (d, $J = 1.5$ Hz, 2H), 3.40 (s, 3H), 3.28 (s, 3H), 3.22 (s, 3H), 3.19 (s, 3H), 3.09 (s, 3H), 2.69 (s, 3H), 2.67 (s, 3H), 2.01 (s, 2H), 1.44–1.25 (m, 8H), 1.06–0.74 (m, 50H); ESI MS m/z 1299 [$\text{C}_{67}\text{H}_{115}\text{N}_{11}\text{O}_{14} + \text{H}$] $^+$.

Example 52 - Preparation of Cyclosporin Alkynyl Ether

[0121] A solution of the acetate of cyclosporin alkynyl ether from Example 51 (61 mg, 0.047 mmol) in methanol (2 mL) was stirred at room temperature. Reaction mixture was treated with potassium carbonate (80 mg, 0.58 mmol) and was allowed to keep stirring under N_2 atmosphere overnight. Mixture was diluted with ethyl acetate, washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford cyclosporin alkynyl ether (31 mg, 52%) as a white solid: ^1H NMR (300 MHz, CDCl_3) δ 7.93 (d, $J = 9.4$ Hz, 1H), 7.63 (d, $J = 7.4$ Hz, 1H), 7.47 (d, $J = 8.1$ Hz, 1H), 7.18 (d, $J = 8.0$ Hz, 1H), 6.19–6.08 (m, 2H), 5.70 (dd, $J = 11.0$, 3.9 Hz, 1H), 5.50 (d, $J = 5.9$ Hz, 1H), 5.45 (d, $J = 15.8$ Hz, 1H), 5.33–5.27 (m, 2H), 5.12 (s, 2H), 5.09–5.01 (m, 1H), 4.93 (dd, $J = 9.8$, 5.8 Hz, 1H), 4.82 (t, $J = 7.4$ Hz, 2H), 4.76–4.61 (m, 2H), 4.53 (t, $J = 7.2$ Hz, 2H), 4.26 (d, $J = 1.7$ Hz, 1H), 4.20 (d, $J = 1.7$ Hz, 3H), 3.78 (t, $J = 6.2$ Hz, 2H), 3.51 (s, 3H), 3.39 (s, 6H), 3.38 (s, 3H), 3.25 (s, 3H), 3.11 (s, 3H), 3.10 (s, 3H), 2.70 (d, $J = 6.0$ Hz, 6H), 1.54 (d, $J = 9.7$ Hz, 2H), 1.46 (s, 1H), 1.36–1.24 (m, 6H), 1.08–0.82 (m, 48H), 0.70 (d, $J = 6.2$ Hz, 2H); ESI MS m/z 1257 [$\text{C}_{65}\text{H}_{113}\text{N}_{11}\text{O}_{13} + \text{H}$] $^+$; HPLC > 99% (AUC), $t_R = 18.99$ min.

Example 53 - Preparation of the Acetate of Cyclosporin Alkynyl Sulfide

[0122] A mixture of acetyl cyclosporin vinyl iodide from Example 3 (70 mg, 0.051 mmol), CuI (5 mg, 0.026 mmol), and $\text{PdCl}_2(\text{PPh}_3)_2$ (18 mg, 0.026 mmol) in

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Et₃N (3 mL) was stirred at room temperature under N₂ atmosphere. Reaction mixture was then treated with propargyl ethyl sulfide (52 mg, 0.51 mmol). Reaction was allowed to stir overnight at room temperature. Solution was filtered through micro-filter and then concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford the acetate of cyclosporin alkynyl sulfide (59 mg, 87%) as an off-white solid: ¹H NMR (300 MHz, CDCl₃) δ 8.43 (d, *J* = 8.6 Hz, 1H), 8.03 (d, *J* = 6.4 Hz, 1H), 7.65 (d, *J* = 8.9 Hz, 1H), 7.57 (d, *J* = 7.9 Hz, 1H), 5.98–5.67 (m, 4H), 5.51 (d, *J* = 10.8 Hz, 2H), 5.36–4.76 (m, 16H), 4.63 (d, *J* = 14.2 Hz, 2H), 4.46 (t, *J* = 7.0 Hz, 2H), 3.98 (s, 2H), 3.43 (s, 3H), 3.25 (s, 3H), 3.22 (s, 3H), 3.19 (s, 3H), 3.15 (s, 3H), 2.68 (s, 3H), 2.67 (s, 3H), 2.01 (s, 2H), 1.34–1.25 (m, 8H), 1.04 (d, *J* = 6.6 Hz, 4H), 1.01–0.79 (m, 50H); ESI MS *m/z* 1329 [C₆₈H₁₁₇N₁₁O₁₃S + H]⁺.

Example 54 - Preparation of Cyclosporin Alkynyl Sulfide

[0123] A solution of the acetate of cyclosporin alkynyl sulfide from Example 53 (59 mg, 0.044 mmol) in methanol (2 mL) was stirred at room temperature. Reaction mixture was treated with potassium carbonate (70 mg, 0.51 mmol) and was allowed to keep stirring under N₂ atmosphere overnight. Mixture was diluted with ethyl acetate, washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford cyclosporin alkynyl sulfide (25 mg, 45%) as a white solid: ¹H NMR (300 MHz, CDCl₃) δ 7.92 (d, *J* = 9.6 Hz, 1H), 7.63 (d, *J* = 7.5 Hz, 1H), 7.46 (d, *J* = 8.4 Hz, 1H), 7.16 (d, *J* = 8.0 Hz, 1H), 6.11–6.02 (m, 2H), 5.70 (dd, *J* = 11.4, 3.9 Hz, 1H), 5.51 (d, *J* = 5.7 Hz, 2H), 5.46–4.91 (m, 16H), 4.82 (t, *J* = 7.5 Hz, 2H), 4.75 (s, 1H), 4.72–4.61 (m, 4H), 4.53 (t, *J* = 7.2 Hz, 2H), 4.26 (d, *J* = 1.7 Hz, 1H), 4.13 (s, 1H), 3.76 (t, *J* = 7.1 Hz, 2H), 3.51 (s, 3H), 3.38 (s, 3H), 3.25 (s, 3H), 3.10 (s, 3H), 3.09 (s, 3H), 2.71 (s, 3H), 2.69 (s, 3H), 1.54 (d, *J* = 9.6 Hz, 1H), 1.45–1.24 (m, 6H), 1.08–0.82 (m, 48H), 0.69 (d, *J* = 6.0 Hz, 1H); ESI MS *m/z* 1287 [C₆₆H₁₁₅N₁₁O₁₂ + H]⁺; HPLC 93.1% (AUC), *t_R* = 20.29 min.

Example 55 - Preparation of the Acetate of Cyclosporin Alkynyl Ether

[0124] A mixture of acetyl cyclosporin vinyl iodide from Example 3 (70 mg, 0.051 mmol), CuI (5 mg, 0.026 mmol), and PdCl₂(PPh₃)₂ (18 mg, 0.026 mmol) in

Et₃N (3 mL) was stirred at room temperature under N₂ atmosphere. Reaction mixture was then treated with propargyl ether (53 µL, 0.51 mmol). Reaction was allowed to stir overnight at room temperature. Solution was filtered through micro-filter and then concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford the acetate of cyclosporin alkynyl ether (28 mg, 41%) as an off-white solid: ¹H NMR (300 MHz, CDCl₃) δ 8.47 (d, *J* = 9.6 Hz, 1H), 8.01 (d, *J* = 6.8 Hz, 1H), 7.51 (d, *J* = 7.1 Hz, 2H), 5.68 (dd, *J* = 10.8, 3.6 Hz, 1H), 5.52 (d, *J* = 8.9 Hz, 2H), 5.38–4.74 (m, 18H), 4.64 (d, *J* = 13.8 Hz, 2H), 4.37 (d, *J* = 1.5 Hz, 3H), 4.25 (d, *J* = 3.6 Hz, 3H), 4.22 (d, *J* = 3.6 Hz, 1H), 3.44 (s, 3H), 3.26 (s, 3H), 3.24 (s, 3H), 3.20 (s, 3H), 3.08 (s, 3H), 2.67 (s, 3H), 2.65 (s, 3H), 2.01 (s, 2H), 1.31 (d, *J* = 7.1 Hz, 4H), 1.27 (d, *J* = 7.3 Hz, 4H), 1.06–0.84 (m, 48H), 0.80 (d, *J* = 6.6 Hz, 2H); ESI MS *m/z* 1323 [C₆₉H₁₁₅N₁₁O₁₄ + H]⁺.

Example 56 - Preparation of Cyclosporin Alkynyl Ether

[0125] A solution of the acetate of cyclosporin alkynyl ether from Example 55 (28 mg, 0.021 mmol) in methanol (2 mL) was stirred at room temperature. Reaction mixture was treated with potassium carbonate (35 mg, 0.25 mmol) and was allowed to keep stirring under N₂ atmosphere overnight. Mixture was diluted with ethyl acetate, washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford cyclosporin alkynyl ether (14 mg, 53%) as a white solid: ¹H NMR (300 MHz, CDCl₃) δ 7.94 (d, *J* = 9.6 Hz, 1H), 7.63 (d, *J* = 7.5 Hz, 1H), 7.45 (d, *J* = 8.2 Hz, 1H), 7.16 (d, *J* = 7.9 Hz, 1H), 6.18–6.09 (m, 2H), 5.70 (dd, *J* = 11.2, 4.1 Hz, 1H), 5.50 (d, *J* = 5.9 Hz, 1H), 5.44 (d, *J* = 15.9 Hz, 1H), 5.29 (dd, *J* = 11.3, 4.3 Hz, 2H), 5.12–4.61 (m, 18H), 4.53 (d, *J* = 7.3 Hz, 1H), 4.37 (d, *J* = 1.7 Hz, 3H), 4.26 (d, *J* = 2.4 Hz, 3H), 3.79 (d, *J* = 6.5 Hz, 1H), 3.58 (s, 3H), 3.51 (s, 3H), 3.26 (s, 3H), 3.11 (s, 3H), 3.10 (s, 3H), 2.70 (s, 3H), 2.68 (s, 3H), 2.43 (t, *J* = 2.9 Hz, 2H), 1.34 (d, *J* = 7.2 Hz, 2H), 1.25 (d, *J* = 6.7 Hz, 2H), 1.08–0.82 (m, 48H), 0.71 (d, *J* = 6.2 Hz, 1H); ESI MS *m/z* 1281 [C₆₇H₁₁₃N₁₁O₁₃ + H]⁺; HPLC >99% (AUC), *t_R* = 18.88 min.

Example 57 - Preparation of the Acetate of Cyclosporin Alkynyl Ether

[0126] A mixture of acetyl cyclosporin vinyl iodide from Example 3 (70 mg, 0.051 mmol), CuI (5 mg, 0.026 mmol), and PdCl₂(PPh₃)₂ (18 mg, 0.026 mmol) in Et₃N (3 mL) was stirred at room temperature under N₂ atmosphere. Reaction mixture was then treated with phenyl propargyl ether (68 mg, 0.51 mmol). Reaction was allowed to stir overnight at room temperature. Solution was filtered through micro-filter and then concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford the acetate of cyclosporin alkynyl ether (67 mg, 96%) as an off-white solid: ¹H NMR (300 MHz, CDCl₃) δ 8.40 (d, *J* = 8.9 Hz, 1H), 8.02 (d, *J* = 6.6 Hz, 1H), 7.74 (d, *J* = 8.7 Hz, 1H), 7.59 (d, *J* = 5.5 Hz, 1H), 7.01–6.84 (m, 3H), 6.01–5.59 (m, 4H), 5.54–5.05 (m, 14H), 4.97 (d, *J* = 10.7 Hz, 2H), 4.81 (d, *J* = 7.7 Hz, 2H), 4.46 (d, *J* = 6.8 Hz, 2H), 3.44 (s, 3H), 3.24 (s, 3H), 3.20 (s, 3H), 3.19 (s, 3H), 3.12 (s, 3H), 2.69 (s, 3H), 2.67 (s, 3H), 2.07 (s, 1H), 2.02 (s, 2H), 1.93 (s, 2H), 1.34 (d, *J* = 7.2 Hz, 4H), 1.32–1.24 (m, 6H), 1.04 (d, *J* = 6.3 Hz, 2H), 1.01–0.73 (m, 48H); ESI MS *m/z* 1361 [C₇₂H₁₁₇N₁₁O₁₄ + H]⁺.

Example 58 - Preparation of Cyclosporin Alkynyl Ether

[0127] A solution of the acetate of cyclosporin alkynyl ether from Example 57 (67 mg, 0.049 mmol) in methanol (2 mL) was stirred at room temperature. Reaction mixture was treated with potassium carbonate (35 mg, 0.58 mmol) and was allowed to keep stirring under N₂ atmosphere overnight. Mixture was diluted with ethyl acetate, washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford cyclosporin alkynyl ether (36 mg, 55%) as a white solid: ¹H NMR (300 MHz, CDCl₃) δ 7.90 (d, *J* = 9.5 Hz, 1H), 7.61 (d, *J* = 7.2 Hz, 1H), 7.43 (d, *J* = 7.8 Hz, 1H), 7.16 (d, *J* = 7.6 Hz, 1H), 6.98 (d, *J* = 7.3 Hz, 2H), 6.20–6.10 (m, 2H), 5.70 (dd, *J* = 10.9, 3.7 Hz, 1H), 5.49 (d, *J* = 5.9 Hz, 1H), 5.44 (d, *J* = 16.1 Hz, 1H), 5.32–4.63 (m, 25H), 4.52 (t, *J* = 7.2 Hz, 2H), 3.50 (s, 3H), 3.38 (s, 3H), 3.25 (s, 3H), 3.10 (s, 6H), 2.70 (s, 3H), 2.68 (s, 3H), 1.40 (d, *J* = 5.7 Hz, 2H), 1.36–0.80 (m, 52H), 0.70 (d, *J* = 6.3 Hz, 2H); ESI MS *m/z* 1319 [C₇₀H₁₁₅N₁₁O₁₃ + H]⁺; HPLC 94.2% (AUC), *t_R* = 20.37 min.

Example 59 - Preparation of the Acetate of Cyclosporin Alkynyl Ester

[0128] A mixture of acetyl cyclosporin vinyl iodide from Example 3 (70 mg, 0.051 mmol), CuI (5 mg, 0.026 mmol), and PdCl₂(PPh₃)₂ (18 mg, 0.026 mmol) in Et₃N (3 mL) was stirred at room temperature under N₂ atmosphere. Reaction mixture was then treated with *t*-butyl (2-propynyloxy)acetate (50 µL, 0.51 mmol). Reaction was allowed to stir for 6 h at room temperature. Solution was filtered through micro-filter and then concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford the acetate of cyclosporin alkynyl ester (71 mg, 98%) as an off-white solid: ¹H NMR (300 MHz, CDCl₃) δ 8.45 (d, *J* = 9.3 Hz, 1H), 8.03 (d, *J* = 6.7 Hz, 1H), 7.70 (d, *J* = 9.0 Hz, 1H), 7.59 (d, *J* = 8.0 Hz, 1H), 6.02–5.56 (m, 5H), 5.52 (d, *J* = 11.2 Hz, 2H), 5.38–4.70 (m, 12H), 4.63 (d, *J* = 14.0 Hz, 2H), 4.45 (t, *J* = 7.0 Hz, 2H), 4.39 (d, *J* = 1.7 Hz, 3H), 4.07 (s, 2H), 3.98 (s, 2H), 3.53 (s, 3H), 3.43 (s, 3H), 3.25 (s, 3H), 3.22 (s, 3H), 3.19 (s, 3H), 3.09 (s, 3H), 2.68 (s, 3H), 2.67 (s, 3H), 2.01 (s, 3H), 1.48 (s, 4H), 1.43–0.20 (m, 8H), 1.05–0.79 (m, 50H); ESI MS *m/z* 1399 [C₇₂H₁₂₃N₁₁O₁₆ + H]⁺.

Example 60 - Preparation of Cyclosporin Alkynyl Ester

[0129] A solution of the acetate of cyclosporin alkynyl ester from Example 59 (71 mg, 0.05 mmol) in methanol (2 mL) was stirred at room temperature. Reaction mixture was treated with potassium carbonate (85 mg, 0.62 mmol) and was allowed to keep stirring under N₂ atmosphere overnight. Mixture was diluted with ethyl acetate, washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford cyclosporin alkynyl acid (28 mg, 41%) as a white solid: ¹H NMR (300 MHz, CDCl₃) δ 7.93 (d, *J* = 10.2 Hz, 1H), 7.59 (d, *J* = 8.7 Hz, 1H), 7.48 (d, *J* = 8.6 Hz, 1H), 7.28 (hidden by solvent peak, 1H), 6.24–6.10 (m, 2H), 5.90–5.66 (m, 2H), 5.45 (d, *J* = 7.2 Hz, 2H), 5.41 (s, 2H), 5.18–4.97 (m, 7H), 4.83 (t, *J* = 7.0 Hz, 1H), 4.74–4.66 (m, 2H), 5.52–4.39 (m, 6H), 4.35 (s, 2H), 4.26 (d, *J* = 1.7 Hz, 1H), 4.19 (s, 3H), 4.14 (d, *J* = 2.0 Hz, 1H), 3.89 (t, *J* = 6.6 Hz, 1H), 3.48 (s, 3H), 3.38 (s, 3H), 3.23 (s, 3H), 3.11 (s, 3H), 3.10 (s, 3H), 2.69 (s, 6H), 1.54 (d, *J* = 9.6 Hz, 1H), 1.46 (s, 1H), 1.38–1.25 (m, 5H), 1.06–0.74 (m, 49H); ESI MS *m/z* 1301 [C₆₆H₁₁₃N₁₁O₁₅ + H]⁺; HPLC > 99% (AUC), *t*_R = 17.09 min.

Example 61 - Preparation of the Acetate of Cyclosporin Alkynyl Acid

[0130] A mixture of acetyl cyclosporin vinyl iodide from Example 3 (70 mg, 0.051 mmol), CuI (5 mg, 0.026 mmol), and PdCl₂(PPh₃)₂ (18 mg, 0.026 mmol) in Et₃N (3 mL) was stirred at room temperature under N₂ atmosphere. Reaction mixture was then treated with 4-pentynoic acid (51 mg, 0.51 mmol). Reaction was allowed to stir for 4 h at room temperature. Solution was filtered through micro-filter and then concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford the acetate of cyclosporin alkynyl acid (23 mg, 34%) as an off-white solid: ¹H NMR (300 MHz, CDCl₃) δ 8.62 (d, *J* = 7.7 Hz, 1H), 8.41 (d, *J* = 9.4 Hz, 1H), 7.97 (d, *J* = 6.8 Hz, 1H), 7.68 (d, *J* = 7.2 Hz, 1H), 5.68 (dd, *J* = 11.1, 3.8 Hz, 2H), 5.59 (d, *J* = 7.6 Hz, 2H), 5.55 (s, 1H), 5.50 (d, *J* = 6.5 Hz, 2H), 5.46–4.77 (m, 13H), 4.63 (d, *J* = 13.4 Hz, 2H), 4.49 (t, *J* = 7.3 Hz, 1H), 4.42 (t, *J* = 6.9 Hz, 2H), 3.44 (s, 3H), 3.35 (s, 3H), 3.25 (s, 3H), 3.20 (s, 3H), 3.12 (s, 3H), 3.08 (s, 3H), 2.67 (s, 3H), 2.65 (s, 3H), 2.01 (s, 2H), 1.36–1.18 (m, 10H), 1.06–0.78 (m, 50H); ESI MS *m/z* 1327 [C₆₈H₁₁₅N₁₁O₁₅ + H]⁺.

Example 62 - Preparation of Cyclosporin Alkynyl Acid

[0131] A solution of the acetate of cyclosporin alkynyl acid from Example 61 (23 mg, 0.017 mmol) in methanol (2 mL) was stirred at room temperature. Reaction mixture was treated with potassium carbonate (30 mg, 0.22 mmol) and was allowed to keep stirring under N₂ atmosphere overnight. Mixture was diluted with ethyl acetate, washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford cyclosporin alkynyl acid (11 mg, 50%) as a white solid: ¹H NMR (300 MHz, CDCl₃) δ 7.94 (d, *J* = 9.4 Hz, 1H), 7.57 (d, *J* = 7.2 Hz, 1H), 7.48 (d, *J* = 8.4 Hz, 1H), 7.28 (hidden by solvent peak, 1H), 6.03–5.66 (m, 3H), 5.46 (d, *J* = 6.1 Hz, 2H), 5.38 (d, *J* = 16.0 Hz, 1H), 5.31 (d, *J* = 8.8 Hz, 1H), 5.10 (d, *J* = 10.8 Hz, 2H), 5.06–4.67 (m, 17H), 4.49 (t, *J* = 7.1 Hz, 2H), 3.81 (t, *J* = 6.5 Hz, 2H), 3.48 (s, 3H), 3.38 (s, 3H), 3.24 (s, 3H), 3.10 (s, 3H), 3.09 (s, 3H), 2.69 (s, 6H), 2.60 (s, 3H), 1.35 (d, *J* = 7.2 Hz, 2H), 1.26 (d, *J* = 6.4 Hz, 2H), 1.07–0.79 (m, 49H), 0.70 (d, *J* = 6.4 Hz, 2H).

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= 6.4 Hz, 2H); ESI MS m/z 1285 [$C_{66}H_{113}N_{11}O_{14} + H$] $^{+}$; HPLC >99% (AUC), t_R = 17.39 min.

Example 63 - Preparation of Cyclosporin Alkynyl Acid

5 [0132] Cyclosporin alkynyl aldehyde from Example 22 (28 mg, 0.02 mmol) was dissolved in acetonitrile (1 mL) and stirred at room temperature under N_2 atmosphere. Reaction was treated with 40% NaH_2PO_4 solution in water (1 mL). Next the mixture was treated with $NaClO_2$ (3 mg, 0.03 mmol) and H_2O_2 (35% in water, 10 3 μ L, 0.03 mmol). Reaction mixture was stirred for 1 h. Reaction was quenched with sodium sulfite (100 mg). Mixture was diluted with ethyl acetate and washed with 1 N HCl. Organic layer was washed with water and brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product (28 mg, 100%) was carried forward without purification. A solution of the crude product (28 mg, 0.021 mmol) in 15 methanol (1.5 mL) was stirred at room temperature. Reaction mixture was treated with potassium carbonate (100 mg, 0.72 mmol) and was allowed to keep stirring under N_2 atmosphere for 4 h. Mixture was diluted with ethyl acetate, washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to 20 afford cyclosporin alkynyl acid (3.5 mg, 13%) as a white solid: 1H NMR (300 MHz, $CDCl_3$) δ 7.89 (d, J = 8.1 Hz, 1H), 7.65 (br s, 1H), 7.39 (br s, 1H), 7.30 (d, J = 7.6 Hz, 1H), 6.33–6.22 (m, 2H), 5.69 (d, J = 10.7 Hz, 2H), 5.60 (d, J = 16.0 Hz, 1H), 5.34 (d, J = 6.8 Hz, 2H), 5.12–4.64 (m, 15H), 4.46 (t, J = 7.0 Hz, 2H), 4.08 (d, J = 6.8 Hz, 2H), 3.50 (s, 3H), 3.39 (s, 3H), 3.23 (s, 3H), 3.15 (s, 3H), 3.09 (s, 3H), 2.69 (s, 3H), 25 2.68 (s, 3H), 1.38 (d, J = 7.3 Hz, 3H), 1.32 (d, J = 7.3 Hz, 3H), 1.27–0.81 (m, 52H); ESI MS m/z 1257 [$C_{64}H_{109}N_{11}O_{14} + H$] $^{+}$; HPLC 83.4% (AUC), t_R = 17.48 min.

Example 64 – Preparation of the Acetate of *cis*-Cyclosporin Vinyl Iodide

30 [0133] To a vigorously stirred suspension of (iodomethyl)triphenylphosphonium iodide (1.3 g, 2.4 mmol) in dry THF (18 mL) under nitrogen, was added sodium bis(trimethylsilyl)amide (2.4 mL, 1 M in THF, 2.4 mmol). After 10 min at room temperature, the mixture was cooled to 0°C and acetyl cyclosporin aldehyde from Example 2 (300 mg, 0.240 mmol) in anhydrous

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THF (10 mL) was added dropwise. After 10 min at 0°C, the reaction was quenched with a saturated solution of ammonium chloride (10 mL), and then allowed to warm to room temperature. The resulting solid was filtered off through a plug of diatomaceous earth and washed with ethyl acetate (200 mL). The organic layer was washed with an aqueous solution of sodium hydrogensulfite (20%, 200 mL), then dried over anhydrous sodium sulfate and concentrated under vacuum to afford crude product (540 mg). The material was purified by semi-preparative HPLC to afford the acetate of the *cis*-isomer of cyclosporin vinyl iodide (150 mg, 46%) as a pale-brown oil: ¹H NMR (300 MHz, CDCl₃) δ 8.53 (d, *J* = 9.8 Hz, 1H), 8.15 (d, *J* = 6.5 Hz, 1H), 7.82 (d, *J* = 9.0 Hz, 1H), 7.63 (d, *J* = 8.2 Hz, 1H), 6.10 (d, *J* = 7.4 Hz, 1H), 6.02–5.94 (m, 1H), 5.69 (dd, *J* = 10.9, 3.8 Hz, 1H), 5.61–5.48 (m, 2H), 5.38–5.13 (m, 3H), 4.98 (d, *J* = 10.9 Hz, 2H), 4.87 (t, *J* = 7.4 Hz, 1H), 4.78 (t, *J* = 9.6 Hz, 1H), 4.63 (d, *J* = 14.2 Hz, 1H), 4.47 (t, *J* = 7.0 Hz, 1H), 3.98 (s, 3H), 3.43 (s, 3H), 3.27 (s, 3H), 3.21 (s, 3H), 3.19 (s, 3H), 3.14 (s, 3H), 2.69 (s, 3H), 2.42–2.30 (m, 1H), 2.22–1.85 (m, 8H), 2.06 (s, 3H), 1.77–1.60 (m, 3H), 1.54–0.75 (m, 56H); ESI MS *m/z* 1357 [C₆₃H₁₁₀IN₁₁O₁₃ + H]⁺.

Example 65 - Preparation of the Acetate of *cis*-Cyclosporin Alkyne

[0134] A mixture of the acetate of *cis*-cyclosporin vinyl iodide from Example 64 (100 mg, 0.073 mmol), CuI (7 mg, 0.037 mmol), and PdCl₂(PPh₃)₂ (26 mg, 0.037 mmol) in Et₃N (3 mL) was stirred at room temperature under N₂ atmosphere. Reaction mixture was then treated with (trimethylsilyl)acetylene (72 mg, 0.737 mmol). Reaction was stirred overnight at room temperature. Solution was filtered through a short pad of silica gel and then filtrate was concentrated under reduced pressure. The crude product was purified by semi-preparative HPLC to afford the acetate of *cis*-cyclosporin alkyne (27 mg, 28%) as an off-white solid: ¹H NMR (300 MHz, CDCl₃) δ 8.40 (d, *J* = 9.7 Hz, 1H), 7.86 (d, *J* = 6.0 Hz, 1H), 7.59 (d, *J* = 9.1 Hz, 1H), 7.37 (d, *J* = 7.6 Hz, 1H), 5.60–5.40 (m, 4H), 5.23 (d, *J* = 11.3 Hz, 1H), 5.15–4.95 (m, 3H), 4.85–4.75 (m, 1H), 4.68 (t, *J* = 7.0 Hz, 1H), 4.64 (t, *J* = 9.5 Hz, 1H), 4.48 (d, *J* = 13.9 Hz, 1H), 4.28 (t, *J* = 7.0 Hz, 1H), 3.27 (s, 3H), 3.10 (s, 3H), 3.06 (s, 3H), 3.01 (s, 3H), 2.95 (s, 3H), 2.52 (s, 3H), 2.51 (s, 3H), 2.30–0.60 (m, 72H), 0.01 (s, 9H); ESI MS *m/z* 1327 [C₆₈H₁₁₉N₁₁O₁₃Si + H].

Example 66 - Preparation of *cis*-Cyclosporin Alkyne

[0135] A solution of acetate of *cis*-cyclosporin alkyne from Example 65
5 (27 mg, 0.020 mmol) in methanol (2 mL) was stirred at room temperature. Reaction mixture was treated with potassium carbonate (12 mg, 0.087 mmol) and allowed to keep stirring under N₂ atmosphere overnight. Mixture was diluted with ethyl acetate, washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-
10 preparative HPLC to afford *cis*-cyclosporin alkyne (12 mg, 48%) as a white solid: ¹H NMR (300 MHz, CDCl₃) δ 8.03 (d, *J* = 9.7 Hz, 1H), 7.64 (d, *J* = 7.2 Hz, 1H), 7.48 (d, *J* = 8.4 Hz, 1H), 7.25 (d, *J* = 7.8 Hz, 1H), 6.10–5.95 (m, 1H), 5.68 (dd, *J* = 11.2, 4.2 Hz, 1H), 5.60–5.50 (m, 1H), 5.45–5.40 (m, 2H), 5.35–5.25 (m, 2H), 5.13–4.95 (m, 2H), 4.85 (t, *J* = 6.9 Hz, 1H), 4.75–4.60 (m, 2H), 4.50 (t, *J* = 7.3 Hz, 1H), 3.93 (t, *J* =
15 6.7 Hz, 1H), 3.52 (s, 3H), 3.40 (s, 3H), 3.24 (s, 3H), 3.12 (s, 3H), 3.11 (s, 3H), 2.69 (s, 6H), 2.50–2.40 (m, 2H), 2.20–0.8 (m, 68H); ESI MS *m/z* 1213 [C₆₃H₁₀₉N₁₁O₁₂ + H]⁺; HPLC >99% (AUC), *t*_R = 18.94 min.

Example 67 - Preparation of the Acetate of *cis*-Cyclosporin Alkyne

20 [0136] To a solution of 1-propynylmagnesium bromide (0.5 M solution in THF, 1.01 mL, 0.55 mmol) in THF (2 mL) was added ZnCl₂ (1.0 M solution in ether, 0.55 mL, 0.55 mmol) dropwise at 0°C. The resulting solution was stirred for 5 min at 0°C. Then ice-bath was removed and reaction mixture was allowed to warm to room
25 temperature. A solution of the acetate of *cis*-cyclosporin vinyl iodide from Example 64 (75 mg, 0.055 mmol) in THF (1 mL) was added followed by the addition of PdCl₂(PPh₃)₂ (19 mg, 0.027 mmol) and the reaction mixture was stirred at room temperature under argon atmosphere overnight. Reaction mixture was concentrated under reduced pressure, diluted with ethyl acetate and filtered through a short pad of
30 silica gel, the filtrate was then concentrated under reduced pressure. The crude product was purified by semi-preparative HPLC to afford the acetate of *cis*-cyclosporin alkyne (47 mg, 67%) as a brown solid: ¹H NMR (500 MHz, CDCl₃) δ 8.56 (d, *J* = 9.7 Hz, 1H), 8.02 (d, *J* = 6.8 Hz, 1H), 7.65 (d, *J* = 9.2 Hz, 1H), 7.52 (d, *J* = 7.7 Hz, 1H), 5.70–5.50 (m, 4H), 5.30–5.10 (m, 3H), 5.00–4.90 (m, 2H), 4.85 (t, *J* =

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7.3 Hz, 1H), 4.76 (t, $J = 9.6$ Hz, 1H), 4.63 (d, $J = 13.9$ Hz, 1H), 4.43 (t, $J = 7.03$ Hz, 1H), 3.41 (s, 3H), 3.27 (s, 3H), 3.22 (s, 3H), 3.20 (s, 3H), 3.10 (s, 3H), 2.68 (s, 3H), 2.67 (s, 3H), 2.45–2.35 (m, 1H), 2.30–0.65 (m, 74H); ESI MS m/z 1269 $[\text{C}_{66}\text{H}_{113}\text{N}_{11}\text{O}_{13} + \text{H}]^+$.

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Example 68 - Preparation of *cis*-Cyclosporin Alkyne

[0137] A solution of the acetate of *cis*-cyclosporin alkyne from Example 67 (37 mg, 0.02 mmol) in methanol (2 mL) was stirred at room temperature. Reaction mixture was treated with potassium carbonate (30 mg, 0.21 mmol) and allowed to keep stirring under N_2 atmosphere overnight. Mixture was diluted with ethyl acetate, washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford *cis*-cyclosporin alkyne (15 mg, 45%) as a white solid: ^1H NMR (300 MHz, CDCl_3) δ 8.09 (d, $J = 9.8$ Hz, 1H), 7.68 (d, $J = 7.2$ Hz, 1H), 7.52 (d, $J = 8.4$ Hz, 1H), 7.30 (d, $J = 7.8$ Hz, 1H), 5.85–5.75 (m, 1H), 5.68 (dd, $J = 10.7$, 3.9 Hz, 1H), 5.43 (d, $J = 7.4$ Hz, 1H), 5.38 (d, $J = 10.0$ Hz, 1H), 5.30 (dd, $J = 11.6$, 4.2 Hz, 1H), 5.15–4.95 (m, 4H), 4.82 (t, $J = 7.2$ Hz, 1H), 4.75–4.65 (m, 2H), 4.50 (t, $J = 7.2$ Hz, 1H), 3.94 (t, $J = 6.8$ Hz, 1H), 3.51 (s, 3H), 3.41 (s, 3H), 3.24 (s, 3H), 3.12 (s, 3H), 3.10 (s, 3H), 2.69 (s, 6H), 2.50–0.81 (m, 72H); ESI MS m/z 1227 $[\text{C}_{64}\text{H}_{111}\text{N}_{11}\text{O}_{12} + \text{H}]^+$; HPLC 98.7% (AUC), $t_R = 19.90$ min.

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Example 69 - Preparation of the Acetates of *cis*- and *trans*-Deuterated Cyclosporin Vinyl Iodide

[0138] A mixture of acetyl cyclosporin aldehyde from Example 2 (500 mg, 0.40 mmol) and iodoform-*d* (1.35 g, 4.0 mmol) in anhydrous THF (10 mL) was cooled to -78°C . After cooling, chromium chloride (1.0 g, 8.0 mmol) was quickly added to the reaction. Mixture was allowed to warm to 0°C and stirred under N_2 atmosphere for 5 h. Mixture was poured into ice-water (300 mL) and extracted with ethyl acetate (3×200 mL). Combined organic layers were washed with brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford the acetate of *trans*-deuterated cyclosporin vinyl iodide (220 mg, 40%) as a light brown solid: ^1H NMR (300 MHz, CDCl_3) δ 8.47 (d,

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$J = 9.6$ Hz, 1H), 8.01 (d, $J = 6.8$ Hz, 1H), 7.57 (t, $J = 8.6$ Hz, 2H), 6.44 (dd, $J = 8.6$, 6.1 Hz, 2H), 6.01–5.56 (m, 4H), 5.52 (d, $J = 10.3$ Hz, 2H), 5.28 (d, $J = 3.4$ Hz, 1H), 5.24 (d, $J = 3.4$ Hz, 1H), 4.97 (d, $J = 10.9$ Hz, 3H), 4.85–4.76 (m, 5H), 4.64 (d, $J = 13.9$ Hz, 2H), 4.43 (t, $J = 7.0$ Hz, 2H), 3.43 (s, 3H), 3.25 (s, 3H), 3.24 (s, 3H), 3.20 (s, 3H), 3.11 (s, 3H), 2.67 (s, 3H), 2.66 (s, 3H), 2.01 (s, 2H), 1.32 (d, $J = 7.1$ Hz, 4H), 1.28 (d, $J = 6.9$ Hz, 4H), 1.06–0.74 (m, 52H); ESI MS m/z 1357 [$C_{63}H_{109}DIN_{11}O_{13} + H$]⁺; and the acetate of *cis*-deuterated cyclosporin vinyl iodide (40 mg, 7%) as a light brown solid: ¹H NMR (300 MHz, CDCl₃) δ 8.57 (d, $J = 9.6$ Hz, 1H), 8.02 (d, $J = 6.8$ Hz, 1H), 7.66 (d, $J = 9.0$ Hz, 1H), 7.53 (d, $J = 7.7$ Hz, 1H), 6.02–5.92 (m, 2H), 5.69 (dd, $J = 11.0$, 3.9 Hz, 1H), 5.54 (d, $J = 3.9$ Hz, 3H), 5.33–5.13 (m, 5H), 4.98 (d, $J = 11.1$ Hz, 3H), 4.82 (t, $J = 7.3$ Hz, 2H), 4.74 (t, $J = 9.5$ Hz, 2H), 4.64 (d, $J = 13.8$ Hz, 2H), 4.32 (t, $J = 7.0$ Hz, 2H), 3.44 (s, 3H), 3.28 (s, 3H), 3.25 (s, 3H), 3.20 (s, 3H), 3.11 (s, 3H), 2.67 (s, 3H), 2.66 (s, 3H), 2.05 (s, 2H), 1.29 (d, $J = 5.5$ Hz, 4H), 1.24 (d, $J = 11.9$ Hz, 4H), 1.05 (d, $J = 6.4$ Hz, 2H), 1.02–0.64 (m, 50H); ESI MS m/z 1357 [$C_{63}H_{109}DIN_{11}O_{13} + H$]⁺.

Example 70 - Preparation of the Acetate of *trans*-Deuterated Cyclosporin Alkyne

[0139] A mixture of the acetate of *trans*-deuterated cyclosporin vinyl iodide from Example 69 (75 mg, 0.055 mmol), CuI (5 mg, 0.028 mmol), and PdCl₂(PPh₃)₂ (19 mg, 0.028 mmol) in Et₃N (3 mL) was stirred at room temperature under N₂ atmosphere. Reaction mixture was then treated with (trimethylsilyl)acetylene (54 mg, 0.55 mmol). Reaction was allowed to stir for 4 h at room temperature. Solution was filtered through micro-filter and then concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford the acetate of *trans*-deuterated cyclosporin alkyne (45 mg, 61%) as an off-white solid: ¹H NMR (300 MHz, CDCl₃) δ 8.21 (d, $J = 9.2$ Hz, 1H), 7.88 (d, $J = 6.6$ Hz, 1H), 7.71 (d, $J = 9.0$ Hz, 1H), 7.47 (d, $J = 7.6$ Hz, 1H), 5.80 (t, $J = 7.3$ Hz, 1H), 5.56–5.35 (m, 3H), 5.29 (d, $J = 10.8$ Hz, 2H), 5.17–4.97 (m, 3H), 4.88 (d, $J = 10.3$ Hz, 3H), 4.86–4.45 (m, 5H), 4.31 (t, $J = 6.8$ Hz, 2H), 3.91 (d, $J = 16.3$ Hz, 1H), 3.80 (s, 3H), 3.25 (s, 3H), 3.08 (s, 3H), 3.05 (s, 3H), 3.01 (s, 3H), 2.93 (s, 3H), 2.53 (s, 3H), 2.51 (s, 3H), 1.85 (s, 1H), 1.81 (s, 2H), 1.21–1.08 (m, 8H), 0.89–0.54 (m, 50H), 0.10 (s, 9H); ESI MS m/z 1328 [$C_{68}H_{118}DN_{11}O_{13}Si + H$]⁺.

Example 71 - Preparation of *trans*-Deuterated Cyclosporin Alkyne

[0140] A solution of acetate of *trans*-deuterated cyclosporin alkyne from
5 Example 70 (44 mg, 0.033 mmol) in methanol (2 mL) was stirred at room
temperature. Reaction mixture was treated with potassium carbonate (55 mg,
0.40 mmol) and was allowed to keep stirring under N₂ atmosphere overnight. Mixture
was diluted with ethyl acetate, washed with saturated sodium bicarbonate solution and
brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was
10 purified by semi-preparative HPLC to afford *trans*-deuterated cyclosporin alkyne
(29 mg, 72%) as a white solid: ¹H NMR (300 MHz, CDCl₃) δ 7.93 (d, *J* = 9.8 Hz,
1H), 7.64 (d, *J* = 7.3 Hz, 1H), 7.46 (d, *J* = 8.3 Hz, 1H), 7.19 (d, *J* = 7.8 Hz, 1H), 5.99
(t, *J* = 7.7 Hz, 2H), 5.70 (dd, *J* = 11.1, 4.1 Hz, 1H), 5.49 (d, *J* = 6.0 Hz, 2H), 5.31 (dd,
J = 10.9, 3.7 Hz, 1H), 5.13–4.62 (m, 16H), 4.52 (t, *J* = 7.3 Hz, 2H), 3.79 (t, *J* =
15 6.5 Hz, 2H), 3.50 (s, 3H), 3.38 (s, 3H), 3.25 (s, 3H), 3.10 (s, 6H), 2.70 (s, 3H), 2.69 (s,
3H), 1.91 (s, 2H), 1.35 (d, *J* = 7.2 Hz, 2H), 1.26 (d, *J* = 5.0 Hz, 2H), 1.08–0.81 (m,
50H), 0.71 (d, *J* = 6.1 Hz, 1H); ESI MS *m/z* 1214 [C₆₃H₁₀₈DN₁₁O₁₂ + H]⁺; HPLC
>99% (AUC), *t*_R = 18.82 min.

Example 72 - Preparation of the Acetate of *trans*-Deuterated Cyclosporin Alkyne

[0141] ZnCl₂ (0.6 mL, 1 M in THF, 0.6 mmol) was added to a solution of
1-propynylmagnesium bromide (1.2 mL, 0.5 M in THF, 0.6 mmol) in THF (3 mL) at
0°C. Mixture was stirred under N₂ atmosphere for 5 min. Mixture was warmed to
25 room temperature and then treated with the acetate of *trans*-deuterated cyclosporin
vinyl iodide from Example 69 (80 mg, 0.060 mmol) and PdCl₂(PPh₃)₂ (20 mg,
0.030 mmol). Mixture stirred for 4 h under N₂ atmosphere at room temperature.
Solution was filtered through micro-filter and then concentrated *in vacuo*. The crude
product was purified by semi-preparative HPLC to afford the acetate of *trans*-
30 deuterated cyclosporin alkyne (62 mg, 83%) as an off-white solid: ¹H NMR
(300 MHz, CDCl₃) δ 8.44 (d, *J* = 9.3 Hz, 1H), 8.04 (d, *J* = 6.7 Hz, 1H), 7.98 (d, *J* =
9.2 Hz, 1H), 7.64 (d, *J* = 7.7 Hz, 1H), 5.83–5.43 (m, 7H), 5.28 (d, *J* = 3.3 Hz, 1H),
5.24 (d, *J* = 3.3 Hz, 1H), 5.18 (d, *J* = 5.7 Hz, 2H), 5.06–4.70 (m, 10H), 4.64 (d, *J* =
14.2 Hz, 1H), 4.46 (t, *J* = 6.9 Hz, 2H), 3.42 (s, 3H), 3.24 (s, 3H), 3.22 (s, 3H), 3.19 (s,

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3H), 3.11 (s, 3H), 2.69 (s, 3H), 2.68 (s, 3H), 2.01 (s, 2H), 1.91 (s, 1H), 1.31 (dd, $J = 10.8, 7.2$ Hz, 6H), 1.04 (d, $J = 6.5$ Hz, 2H), 1.01–0.70 (m, 52H); ESI MS m/z 1270 $[C_{66}H_{112}DN_{11}O_{13} + H]^+$.

5 **Example 73 – Preparation of *trans*-Deuterated Cyclosporin Alkyne**

[0142] A solution of the acetate of *trans*-deuterated cyclosporine alkyne from Example 72 (62 mg, 0.049 mmol) in methanol (2 mL) was stirred at room temperature. Reaction mixture was treated with potassium carbonate (70 mg, 10 0.50 mmol) and was allowed to keep stirring under N₂ atmosphere overnight. Mixture was diluted with ethyl acetate, washed with saturated sodium bicarbonate solution and brine, dried over sodium sulfate, and concentrated *in vacuo*. The crude product was purified by semi-preparative HPLC to afford *trans*-deuterated cyclosporine alkyne (31 mg, 52%) as a white solid: ¹H NMR (300 MHz, CDCl₃) δ 7.92 (d, $J = 9.7$ Hz, 15 1H), 7.63 (d, $J = 7.3$ Hz, 1H), 7.44 (d, $J = 8.2$ Hz, 1H), 7.20 (d, $J = 7.8$ Hz, 1H), 6.23 (t, $J = 7.3$ Hz, 1H), 5.49 (d, $J = 6.0$ Hz, 2H), 5.31 (dd, $J = 10.7, 3.4$ Hz, 1H), 5.11–4.62 (m, 20H), 4.52 (t, $J = 7.2$ Hz, 2H), 3.82 (t, $J = 6.5$ Hz, 2H), 3.51 (s, 3H), 3.38 (s, 3H), 3.25 (s, 3H), 3.11 (s, 3H), 3.10 (s, 3H), 2.70 (s, 3H), 2.68 (s, 3H), 1.34 (d, $J = 7.2$ Hz, 2H), 1.26 (d, $J = 4.8$ Hz, 2H), 1.08–0.81 (m, 52H), 0.73 (d, $J = 6.4$ Hz, 1H); 20 ESI MS m/z 1228 $[C_{64}H_{110}DN_{11}O_{12} + H]^+$; HPLC >99% (AUC), $t_R = 19.47$ min.

Example 74 – Preparation of Cyclosporin Diol

[0143] To a mechanically stirred solution of diisopropylamine (2.6 mL, 25 18 mmol) in THF (50 mL) at –78°C was added dropwise *n*-butyllithium (6.6 mL, 2.5 M in hexane, 17 mmol), then the mixture was stirred for 0.5 h. A solution of cyclosporin A (1.0 g, 0.83 mmol) in THF (8 mL) was added, and then the mixture was stirred for 2 h at –78°C. Paraformaldehyde (8.0 g) was heated to 170°C and the resulting formaldehyde gas was transferred into the reaction via a glass tube which 30 was wrapped with cotton and aluminum foil over 2 h. After stirred another 1 h at –78°C, the reaction mixture was quenched with water (10 mL). The mixture was allowed to warm to room temperature, diluted with ethyl acetate (150 mL) and washed with water (2 × 50 mL). The organic layer was separated, dried over anhydrous sodium sulfate and concentrated under vacuum. The crude material was

purified by semi-preparative HPLC to afford cyclosporin diol (0.45 g, 44%) as a white solid: ^1H NMR (300 MHz, CDCl_3) δ 8.09 (d, $J = 9.9$ Hz, 1H), 7.70 (d, $J = 7.4$ Hz, 1H), 7.57 (d, $J = 8.2$ Hz, 1H), 7.15 (overlapped with CHCl_3 , 1H), 5.70 (dd, $J = 11.0, 4.0$ Hz, 1H), 5.49 (d, $J = 6.4$ Hz, 1H), 5.38–5.30 (m, 3H), 5.16–4.93 (m, 5H), 4.83 (t, $J = 7.2$ Hz, 1H), 4.65 (t, $J = 9.5$ Hz, 1H), 4.54 (t, $J = 7.2$ Hz, 1H), 4.05 (d, $J = 6.8$ Hz, 2H), 3.73 (t, $J = 6.3$ Hz, 1H), 3.49 (s, 3H), 3.30 (s, 3H), 3.25 (s, 3H), 3.15 (s, 3H), 3.11 (s, 3H), 2.70 (s, 3H), 2.69 (s, 3H), 2.50–2.38 (m, 2H), 2.20–1.92 (m, 6H), 1.75–0.65 (m, 64H); ESI MS m/z 1233 [$\text{C}_{63}\text{H}_{113}\text{N}_{11}\text{O}_{13} + \text{H}$] $^+$.

10 **Example 75 - Preparation of Cyclosporin Diacetate**

[0144] To a solution of cyclosporin diol from Example 74 (0.43 g, 0.35 mmol) in methylene chloride (5 mL) was added pyridine (0.57 mL, 7.0 mmol) followed by 4-(dimethylamino)pyridine (86 mg, 0.70 mmol) and acetic anhydride (1.0 mL, 10.5 mmol). The reaction mixture was stirred for 2 d at room temperature. The reaction was diluted with ethyl ether (150 mL), washed with a saturated solution of sodium bicarbonate (30 mL), 1N HCl solution (30 mL) and brine (30 mL). The organic layer was separated, dried over anhydrous sodium sulfate and concentrated under vacuum. The crude material was purified by semi-preparative HPLC to afford cyclosporin diacetate (0.23 g, 50%) as a white solid: ^1H NMR (300 MHz, CDCl_3) δ 8.60 (d, $J = 9.8$ Hz, 1H), 8.05 (d, $J = 6.6$ Hz, 1H), 7.55 (d, $J = 7.8$ Hz, 1H), 7.49 (d, $J = 9.3$ Hz, 1H), 5.68 (dd, $J = 11.0, 4.0$ Hz, 1H), 5.49 (s, 2H), 5.40–4.95 (m, 8H), 4.85 (t, $J = 7.5$ Hz, 1H), 4.76 (t, $J = 9.3$ Hz, 1H), 4.58–4.34 (m, 3H), 3.37 (s, 3H), 3.27 (s, 3H), 3.23 (s, 3H), 3.20 (s, 3H), 3.14 (s, 3H), 2.67 (s, 3H), 2.66 (s, 3H), 2.48–2.35 (m, 1H), 2.10 (s, 3H), 2.01 (s, 3H), 1.98–1.85 (m, 2H), 1.75–0.65 (m, 67H); ESI MS m/z 1317 [$\text{C}_{67}\text{H}_{117}\text{N}_{11}\text{O}_{15} + \text{H}$] $^+$.

Example 76 - Preparation of Cyclosporin Aldehyde

30 [0145] Ozone was bubbled into a solution of cyclosporin diacetate from Example 75 (0.22 g, 0.17 mmol) in methylene chloride (10 mL) at -78°C until a blue color was developed. The mixture was degassed with nitrogen for a few min and dimethylsulfide (0.4 mL) was added at -78°C . The reaction mixture was allowed to warm to room temperature and stirred for 3 h. The reaction mixture was concentrated

in vacuo and the residue was dissolved in ethyl acetate (120 mL), washed with water (2 × 20 mL) and brine (30 mL), dried over sodium sulfate, filtered, and concentrated *in vacuo* to afford cyclosporin aldehyde (0.19 g, 86%) as a white solid. The crude was carried to the next step without further purification: ¹H NMR (300 MHz, CDCl₃)
5 δ 9.55 (d, *J* = 3.4 Hz, 1H), 8.60 (d, *J* = 9.9 Hz, 1H), 7.96 (d, *J* = 7.1 Hz, 1H), 7.53 (d, *J* = 7.7 Hz, 1H), 7.33 (d, *J* = 9.1 Hz, 1H), 5.68 (dd, *J* = 11.0, 4.0 Hz, 1H), 5.53 (d, *J* = 11.2 Hz, 1H), 5.47 (d, *J* = 11.2 Hz, 1H), 5.30 (dd, *J* = 12.3, 3.6 Hz, 1H), 5.18–4.92 (m, 5H), 4.84 (t, *J* = 6.9 Hz, 1H), 4.72 (t, *J* = 9.6 Hz, 1H), 4.55–4.35 (m, 3H), 3.39 (s, 3H), 3.30 (s, 3H), 3.29 (s, 3H), 3.21 (s, 3H), 3.12 (s, 3H), 2.66 (s, 3H), 2.65 (s, 3H),
10 2.48–2.30 (m, 3H), 2.10 (s, 3H), 1.99 (s, 3H), 1.80–0.75 (m, 64H); ESI MS *m/z* 1305 [C₆₅H₁₁₃N₁₁O₁₆ + H]⁺.

Example 77 - Preparation of the Acetates of *cis*- and *trans*-Cyclosporin Vinyl Iodide

15 [0146] To an ice-cooled suspension of chromium(II) chloride (664 mg, 5.4 mmol) in THF (7 mL) was added a solution of cyclosporin aldehyde from Example 76 (0.35 g, 0.27 mmol) and iodoform (850 mg, 2.16 mmol) in THF (7 mL), then the mixture was stirred at 0°C. After 4 h, the reaction mixture was poured into
20 ice-water (40 mL). The aqueous layer was separated and extracted with ethyl acetate (3 × 50 mL). The combined organics were dried over anhydrous sodium sulfate and concentrated under vacuum. The crude was purified by semi-preparative HPLC to afford the acetate of *cis*-cyclosporin vinyl iodide (26 mg, 7%): ¹H NMR (CDCl₃, 300 MHz) δ 8.62 (d, *J* = 9.6 Hz, 1H), 8.02 (d, *J* = 6.9 Hz, 1H), 7.60 (d, *J* = 7.5 Hz, 1H), 7.55 (d, *J* = 9.3 Hz, 1H), 6.07 (d, *J* = 7.4 Hz, 1H), 6.02–5.92 (m, 1H), 5.69 (dd, *J* = 11.1, 3.9 Hz, 1H), 5.50 (dd, *J* = 15.6, 11.7 Hz, 2H), 5.30–4.93 (m, 6H), 4.85 (t, *J* = 7.2 Hz, 1H), 4.74 (t, *J* = 9.3 Hz, 1H), 4.55–4.32 (m, 3H), 3.36 (s, 3H), 3.29 (s, 3H), 3.23 (s, 3H), 3.20 (s, 3H), 3.14 (s, 3H), 2.67 (s, 6H), 2.43–2.32 (m, 1H), 2.23–2.15 (m, 2H), 2.10 (s, 3H), 2.06 (s, 3H), 1.95–1.82 (m, 3H), 1.75–1.60 (m, 3H), 1.48–0.82
25 (m, 58H); ESI MS *m/z* 1429 [C₆₆H₁₁₄IN₁₁O₁₅ + H]⁺; the acetate of *trans*-cyclosporin vinyl iodide (0.18 g, 47%): ¹H NMR (CDCl₃, 300 MHz) δ 8.54 (d, *J* = 9.9 Hz, 1H), 8.00 (d, *J* = 6.6 Hz, 1H), 7.58 (d, *J* = 7.5 Hz, 1H), 7.43 (d, *J* = 9.0 Hz, 1H), 6.45–6.35 (m, 1H), 5.83 (d, *J* = 14.4 Hz, 1H), 5.69 (dd, *J* = 10.8, 3.9 Hz, 1H), 5.55–5.43 (m,

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2H), 5.38 (dd, $J = 12.0, 3.9$ Hz, 1H), 5.30–4.93 (m, 5H), 4.85 (t, $J = 6.9$ Hz, 1H), 4.77 (t, $J = 9.6$ Hz, 1H), 4.53–4.30 (m, 3H), 3.36 (s, 3H), 3.27 (s, 3H), 3.26 (s, 3H), 3.20 (s, 3H), 3.15 (s, 3H), 2.67 (s, 3H), 2.66 (s, 3H), 2.43–2.32 (m, 1H), 2.23–2.11 (m, 2H), 2.10 (s, 3H), 2.02 (s, 3H), 1.95–1.60 (m, 6H), 1.48–0.75 (m, 58H); ESI MS m/z 1429

5 $[\text{C}_{66}\text{H}_{114}\text{IN}_{11}\text{O}_{15} + \text{H}]^+$.

Example 78 - Preparation of the Acetate of Cyclosporin Alkyne

[0147] To a mixture of the acetate of *trans*-cyclosporin vinyl iodide from

10 Example 77 (60 mg, 0.042 mmol), copper(I) iodide (4 mg, 0.021 mmol), dichlorobis(triphenylphosphine)palladium(II) (15 mg, 0.021 mmol) in triethylamine (2 mL) was added (trimethylsilyl)acetylene (60 μL , 0.42 mmol), then the mixture was stirred at room temperature for 3 h. The reaction mixture was filtered through a micro-filter and concentrated under vacuum. The crude was purified by semi-

15 preparative HPLC to afford the acetate of cyclosporin alkyne (16 mg, 28%) as a white solid: ^1H NMR (CDCl_3 , 300 MHz) δ 8.53 (d, $J = 9.3$ Hz, 1H), 8.06 (d, $J = 6.6$ Hz, 1H), 7.68 (app t, $J = 8.7$ Hz, 2H), 6.02 (dt, $J = 15.9, 7.5$ Hz, 1H), 5.70 (dd, $J = 10.8, 3.9$ Hz, 1H), 5.55–5.37 (m, 2H), 5.35–5.13 (m, 4H), 5.06–4.93 (m, 4H), 4.87 (t, $J = 7.2$ Hz, 1H), 4.75 (t, $J = 9.3$ Hz, 1H), 4.53–4.33 (m, 4H), 3.38 (s, 3H), 3.27 (s, 3H),

20 3.24 (s, 3H), 3.19 (s, 3H), 3.12 (s, 3H), 2.69 (s, 3H), 2.68 (s, 3H), 2.43–2.32 (m, 1H), 2.23–2.11 (m, 2H), 2.11 (s, 3H), 2.01 (s, 3H), 1.95–1.60 (m, 6H), 1.48–0.75 (m, 56H), 0.17 (s, 9H); ESI MS m/z 1399 $[\text{C}_{71}\text{H}_{123}\text{N}_{11}\text{O}_{15}\text{Si} + \text{H}]^+$.

Example 79 - Preparation of Cyclosporin Alkyne

25 [0148] To a solution of the acetate of cyclosporin alkyne from Example 78 (16 mg, 0.011 mmol) in MeOH (2 mL) was added potassium carbonate (46 mg, 0.33 mmol), then the mixture was stirred at room temperature. After 12 h, the reaction mixture was quenched with a saturated solution of ammonium chloride

30 (10 mL). The aqueous layer was extracted with ethyl acetate (3×20 mL). The combined organics were dried over anhydrous sodium sulfate and concentrated under vacuum. The crude material was purified by semi-preparative HPLC to afford cyclosporin alkyne (8 mg, 57%) as a white solid: ^1H NMR (300 MHz, CDCl_3) δ 8.07 (d, $J = 9.9$ Hz, 1H), 7.68 (d, $J = 8.4$ Hz, 1H), 7.45 (d, $J = 8.1$ Hz, 1H), 7.20 (d, $J =$

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7.8 Hz, 1H), 6.28–6.15 (m, 1H), 5.70 (dd, $J = 10.8, 3.6$ Hz, 1H), 5.50 (d, $J = 6.6$ Hz, 1H), 5.39 (d, $J = 15.3$ Hz, 1H), 5.30 (dd, $J = 11.7, 3.6$ Hz, 1H), 5.12–4.92 (m, 5H), 4.84 (t, $J = 6.9$ Hz, 1H), 4.66 (t, $J = 9$ Hz, 1H), 4.51 (t, $J = 7.2$ Hz, 1H), 4.04 (d, $J = 6.6$ Hz, 2H), 3.76 (t, $J = 6.0$ Hz, 1H), 3.51 (s, 3H), 3.31 (s, 3H), 3.27 (s, 3H), 3.15 (s, 3H), 3.11 (s, 3H), 2.69 (s, 3H), 2.68 (s, 3H), 2.55–2.38 (m, 5H), 2.20–1.90 (m, 6H), 1.80–1.60 (m, 5H), 1.45–0.75 (m, 54H); ESI MS m/z 1243 [$C_{64}H_{111}N_{11}O_{13} + H$]⁺; HPLC 96.2% (AUC), $t_R = 13.18$ min.

Example 80 - Preparation of the Acetate of Cyclosporin Alkyne

[0149] To an ice-cooled solution of 1-(trimethylsilyl)-1-propyne- d_3 (49 mg, 0.42 mmol) in triethylamine (2 mL) was added tetrabutylammonium fluoride (0.5 mL, 1 M in THF, 0.5 mmol), then the mixture was stirred for 10 min. The reaction mixture was allowed to warm to room temperature, then the solution was transferred into a mixture of the acetate of trans-cyclosporin vinyl iodide from Example 77 (60 mg, 0.042 mmol), copper(I) iodide (4 mg, 0.021 mmol), dichlorobis(triphenylphosphine)palladium(II) (15 mg, 0.021 mmol). The resulting reaction mixture was stirred at room temperature for 3 h. The reaction mixture was filtered through a micro-filter and concentrated under vacuum. The crude was purified by semi-preparative HPLC to afford the acetate of cyclosporin alkyne (47 mg, 84%) as a white solid: ¹H NMR (CDCl₃, 300 MHz) δ 8.56 (d, $J = 9.9$ Hz, 1H), 8.04 (d, $J = 6.6$ Hz, 1H), 7.68 (d, $J = 7.5$ Hz, 1H), 7.60 (d, $J = 9.3$ Hz, 1H), 5.88–5.77 (m, 1H), 5.70 (dd, $J = 10.8, 3.9$ Hz, 1H), 5.57–5.42 (m, 2H), 5.40 (dd, $J = 12.0, 3.9$ Hz, 1H), 5.30–5.20 (m, 2H), 5.17 (t, $J = 7.5$ Hz, 1H), 5.05–4.94 (m, 3H), 4.90–4.75 (m, 2H), 4.57–4.35 (m, 3H), 3.37 (s, 3H), 3.26 (s, 3H), 3.24 (s, 3H), 3.19 (s, 3H), 3.14 (s, 3H), 2.68 (s, 6H), 2.45–2.37 (m, 1H), 2.30–2.12 (m, 2H), 2.11 (s, 3H), 2.02 (s, 3H), 1.95–1.65 (m, 6H), 1.48–0.75 (m, 58H); ESI MS m/z 1344 [$C_{69}H_{114}D_3N_{11}O_{15} + H$]⁺.

Example 81 - Preparation of Cyclosporin Alkyne

[0150] To a solution of the acetate of cyclosporin alkyne from Example 80 (45 mg, 0.033 mmol) in MeOH (3 mL) was added potassium carbonate (137 mg, 0.99 mmol), then the mixture was stirred at room temperature. After 12 h, the

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reaction mixture was quenched with a saturated solution of ammonium chloride (15 mL). The aqueous layer was extracted with ethyl acetate (3 × 30 mL). The combined organics were dried over anhydrous sodium sulfate and concentrated under vacuum. The crude material was purified by semi-preparative HPLC to afford cyclosporin alkyne (24 mg, 57%) as a white solid: ¹H NMR (300 MHz, CDCl₃) δ 8.07 (d, *J* = 9.6 Hz, 1H), 7.68 (d, *J* = 7.2 Hz, 1H), 7.47 (d, *J* = 8.1 Hz, 1H), 7.20 (d, *J* = 8.1 Hz, 1H), 6.05–5.95 (m, 1H), 5.70 (dd, *J* = 10.8, 3.6 Hz, 1H), 5.50 (d, *J* = 6.6 Hz, 1H), 5.35 (d, *J* = 16.2 Hz, 1H), 5.30 (dd, *J* = 11.7, 3.6 Hz, 1H), 5.12–4.90 (m, 5H), 4.83 (t, *J* = 6.9 Hz, 1H), 4.65 (t, *J* = 9.0 Hz, 1H), 4.54 (t, *J* = 7.2 Hz, 1H), 4.04 (d, *J* = 6.6 Hz, 2H), 3.72 (t, *J* = 6.3 Hz, 1H), 3.50 (s, 3H), 3.30 (s, 3H), 3.27 (s, 3H), 3.14 (s, 3H), 3.11 (s, 3H), 2.70 (s, 3H), 2.69 (s, 3H), 2.55–2.35 (m, 2H), 2.20–1.90 (m, 5H), 1.80–1.58 (m, 6H), 1.45–0.75 (m, 56H); ESI MS *m/z* 1260 [C₆₅H₁₁₀D₃N₁₁O₁₃ + H]⁺; HPLC >99% (AUC), *t*_R = 13.48 min.

Example 82 - Preparation of the Acetate of Cyclosporin Alkyne

[0151] To an ice-cooled solution of 1-propynylmagnesium bromide (0.76 mL, 0.5 M in THF, 0.38 mmol) in THF (1 mL) was added a solution of zinc chloride (0.38 mL, 1 M in ethyl ether, 0.38 mmol). The reaction was stirred at 0°C for 10 min, and then allowed to warm to room temperature. A solution of the acetate of trans-cyclosporin vinyl iodide from Example 77 (54 mg, 0.038 mmol) in THF (2 mL) was added into the reaction mixture followed by copper(I) iodide (2 mg, 0.008 mmol) and dichlorobis(triphenylphosphine)palladium(II) (3 mg, 0.004 mmol). The resulting reaction mixture was stirred at room temperature for 1 h, and then quenched with a saturated solution of ammonium chloride. The aqueous layer was extracted with ethyl acetate (3 × 30 mL). The combined organics were dried over anhydrous sodium sulfate and concentrated under vacuum. The crude was purified by semi-preparative HPLC to afford the acetate of cyclosporin alkyne (27 mg, 53%) as a white solid: ¹H NMR (CDCl₃, 300 MHz) δ 8.55 (d, *J* = 9.6 Hz, 1H), 8.04 (d, *J* = 6.9 Hz, 1H), 7.68 (d, *J* = 7.8 Hz, 1H), 7.60 (d, *J* = 9.0 Hz, 1H), 5.88–5.75 (m, 1H), 5.70 (dd, *J* = 10.8, 3.9 Hz, 1H), 5.53 (d, *J* = 11.4 Hz, 1H), 5.45 (d, *J* = 11.1 Hz, 1H), 5.39 (dd, *J* = 12.0, 3.9 Hz, 1H), 5.30–5.20 (m, 2H), 5.17 (t, *J* = 7.5 Hz, 1H), 5.05–4.94 (m, 3H), 4.90–4.75 (m, 2H), 4.57–4.35 (m, 3H), 3.37 (s, 3H), 3.26 (s, 3H), 3.24 (s, 3H), 3.19 (s, 3H),

3.13 (s, 3H), 2.68 (s, 6H), 2.45–2.35 (m, 1H), 2.30–2.12 (m, 2H), 2.11 (s, 3H), 2.02 (s, 3H), 1.95–1.65 (m, 10H), 1.48–0.75 (m, 57H); ESI MS m/z 1341 [$C_{69}H_{117}N_{11}O_{15} + H$]⁺.

5 **Example 83 - Preparation of Cyclosporin Alkyne**

[0152] To a solution of the acetate of cyclosporin alkyne from Example 82 (25 mg, 0.019 mmol) in MeOH (2 mL) was added potassium carbonate (79 mg, 0.57 mmol), then the mixture was stirred at room temperature. After 8 h, the reaction mixture was quenched with a saturated solution of ammonium chloride (10 mL). The aqueous layer was extracted with ethyl acetate (3 × 20 mL). The combined organics were dried over anhydrous sodium sulfate and concentrated under vacuum. The crude material was purified by semi-preparative HPLC to afford cyclosporin alkyne (14 mg, 58%): ¹H NMR (300 MHz, CDCl₃) δ 8.06 (d, J = 9.9 Hz, 1H), 7.68 (d, J = 7.5 Hz, 1H), 7.45 (d, J = 8.1 Hz, 1H), 7.18 (d, J = 8.1 Hz, 1H), 6.05–5.95 (m, 1H), 5.70 (dd, J = 10.8, 3.6 Hz, 1H), 5.50 (d, J = 6.6 Hz, 1H), 5.40–5.25 (m, 2H), 5.12–4.90 (m, 5H), 4.83 (t, J = 6.9 Hz, 1H), 4.65 (t, J = 9 Hz, 1H), 4.54 (t, J = 7.2 Hz, 1H), 4.04 (d, J = 6.6 Hz, 2H), 3.73 (t, J = 6.3 Hz, 1H), 3.50 (s, 3H), 3.30 (s, 3H), 3.27 (s, 3H), 3.14 (s, 3H), 3.11 (s, 3H), 2.70 (s, 3H), 2.68 (s, 3H), 2.55–2.35 (m, 2H), 2.20–1.98 (m, 5H), 1.92 (d, J = 1.7 Hz, 3H), 1.80–1.55 (m, 6H), 1.48–0.70 (m, 56H); ESI MS m/z 1257 [$C_{65}H_{113}N_{11}O_{13} + H$]⁺; HPLC >99% (AUC), t_R = 13.41 min.

Example 84 - Preparation of the Acetate of Cyclosporin Alkyne

25 [0153] To a dried 25 mL flask charged with a solution of the acetate of *trans*-cyclosporin vinyl iodide from Example 77 (60 mg, 0.04 mmol) in triethylamine (2 mL) was added copper(I) iodide (3.8 mg, 0.02 mmol), followed by dichlorobis(triphenylphosphine)palladium(II) (7.0 mg, 0.01 mmol), and propargyl alcohol (0.02 mL, 0.40 mmol). After 2.5 h, the reaction mixture was filtered through a microfilter, and then concentrated under reduced pressure. The crude material was purified by semi-preparative HPLC to yield the acetate of cyclosporin alkyne (54 mg, quantitative): ¹H NMR (300 MHz, CDCl₃) δ 8.47 (d, J = 9.8 Hz, 1H), 7.94 (d, J = 6.8 Hz, 1H), 7.42 (d, J = 7.7 Hz, 1H), 7.27 (d, J = 7.6 Hz, 1H), 5.38–5.26 (m, 1H), 5.47 (s, 2H), 5.45–5.32 (m, 1H), 5.22 (dd, J = 9.3, 3.2 Hz, 1H), 5.15 (dd, J = 7.7,

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5.8 Hz, 1H), 5.07–4.91 (m, 2H), 4.83 (t, $J = 6.9$ Hz, 1H), 4.55–4.21 (m, 4H), 3.35 (s, 3H), 3.27 (s, 3H), 3.26 (s, 3H), 3.21 (s, 3H), 3.15 (s, 3H), 2.67 (s, 3H), 2.65 (s, 3H), 2.44–2.11 (m, 1H), 2.10 (s, 3H), 2.03 (s, 3H), 2.01–1.59 (m, 4H), 1.49–0.74 (m, 68H); ESI MS m/z 1357 [$C_{69}H_{117}N_{11}O_{16} + H$]⁺.

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Example 85 - Preparation of Cyclosporin Alkyne

[0154] To a dried 25 mL flask charged with a solution of the acetate of cyclosporin alkyne from Example 84 (54 mg, 0.04 mmol) in methanol (2 mL) was added potassium carbonate (110 mg, 0.80 mmol). This was allowed to stir at room temperature for 5 h, after which it was quenched with a saturated solution of ammonium chloride (8 mL). This was allowed to stir for 10 min. The aqueous layer was then extracted with ethyl acetate (3 × 40 mL). The combined organics were washed with brine (15 mL), dried over anhydrous sodium sulfate, and then concentrated under reduced pressure. The crude material was purified by semi-preparative HPLC to yield cyclosporin alkyne (28 mg, 54%) as a white solid: ¹H NMR (300 MHz, CDCl₃) δ 7.80 (d, $J = 9.7$ Hz, 1H), 7.55–7.47 (m, 2H), 7.32 (d, $J = 7.7$ Hz, 1H), 6.05–5.92 (m, 1H), 5.67 (dd, $J = 10.7, 3.5$ Hz, 1H), 5.48–5.35 (m, 3H), 5.14–4.86 (m, 6H), 4.81 (t, $J = 7.2$ Hz, 1H), 4.74 (t, $J = 9.3$ Hz, 1H), 4.47 (t, $J = 7.1$ Hz, 1H), 4.34 (d, $J = 5.2$ Hz, 1H), 4.09–3.96 (m, 3H), 3.49 (s, 3H), 3.31 (s, 3H), 3.24 (s, 3H), 3.18 (s, 3H), 3.14 (s, 3H), 3.10 (s, 3H), 2.69 (s, 6H), 2.48–2.37 (m, 1H), 2.32–2.03 (m, 5H), 2.01–0.77 (m, 61H); ESI MS m/z 1274 [$C_{65}H_{113}N_{11}O_{14} + H$]⁺; HPLC >99% (AUC), $t_R = 17.32$ min.

Example 86 - Preparation of the Acetate of Cyclosporin Alkyne Alcohol

[0155] To a dried 25 mL flask charged with a solution of the acetate of *trans*-cyclosporin vinyl iodide from Example 77 (120 mg, 0.08 mmol) in triethylamine (3 mL) was added copper(I) iodide (15 mg, 0.08 mmol) followed by dichlorobis(triphenylphosphine)palladium(II) (21 mg, 0.04 mmol) and 3-butyne-2-ol (0.06 mL, 0.80 mmol). After 2.5 h, the reaction mixture was filtered through a microfilter then concentrated under reduced pressure. The crude material was purified by semi-preparative HPLC to yield the acetate of cyclosporin alkyne alcohol (79 mg, 73%): ¹H NMR (300 MHz, CDCl₃) δ 8.49 (d, $J = 4.7$ Hz, 0.5H), 8.46 (d, $J =$

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4.6 Hz, 0.5H), 7.98 (d, $J = 6.0$ Hz, 0.5H), 7.96 (d, $J = 5.9$ Hz, 0.5H), 7.49 (d, $J = 7.0$ Hz, 0.5H), 7.46 (d, $J = 6.7$ Hz, 0.5H), 7.38 (d, $J = 9.1$ Hz, 0.5H), 7.34 (d, $J = 9.1$ Hz, 0.5H), 5.79–5.58 (m, 1H), 5.47 (s, 2H), 5.43–5.09 (m, 3H), 5.04–4.78 (m, 4H), 4.76–4.33 (m, 3H), 3.37 (s, 1.5H), 3.35 (s, 1.5H), 3.25 (s, 6H), 3.20 (s, 3H), 3.14 (s, 1.5H), 3.13 (s, 1.5H), 2.67 (s, 3H), 2.66 (s, 3H), 2.49–2.18 (m, 1H), 2.10 (s, 3H), 2.03 (s, 3H), 1.98–1.58 (m, 4H), 1.47–0.71 (m, 70H); ESI MS m/z 1371 $[\text{C}_{70}\text{H}_{119}\text{N}_{11}\text{O}_{16} + \text{H}]^+$.

Example 87 - Preparation of Cyclosporin Alkyne

[0156] To a dried 25 mL flask charged with a solution of the acetate of cyclosporin alkyne from Example 86 (25 mg, 0.018 mmol) in methanol (1.5 mL) was added potassium carbonate (50 mg, 0.36 mmol). This was allowed to stir at room temperature for 3.5 h, after which it was quenched with a saturated solution of ammonium chloride (1.5 mL). This was allowed to stir for 5 min. The aqueous layer was then extracted with ethyl acetate (2×20 mL). The combined organics were washed with brine (15 mL), dried over anhydrous sodium sulfate, and then concentrated under reduced pressure. The crude material was purified twice by semi-preparative HPLC to yield cyclosporin alkyne (11 mg, 48%) as a white solid and mixture of diastereomers: ^1H NMR (300 MHz, CDCl_3) δ 7.89 (d, $J = 10.1$ Hz, 0.7H), 7.70 (d, $J = 10.3$ Hz, 0.3H), 7.54–7.44 (m, 2H), 7.30–7.21 (m, 1H), 6.02–5.86 (m, 1H), 5.67 (dd, $J = 7.1, 3.9$ Hz, 1H), 5.49–5.28 (m, 3H), 5.17–4.42 (m, 9H), 4.09–3.91 (m, 3H), 3.50 (s, 3H), 3.31 (s, 3H), 3.24 (s, 3H), 3.14 (s, 3H), 3.10 (s, 2.1H), 3.09 (s, 0.9H), 2.68 (s, 6H), 2.49–2.30 (m, 1H), 2.29–1.89 (m, 5H), 1.84–0.68 (m, 67H); ESI MS m/z 1287 $[\text{C}_{66}\text{H}_{115}\text{N}_{11}\text{O}_{14} + \text{H}]^+$; HPLC >99% (AUC), $t_R = 17.96$ min.

Example 88 - Preparation of the Diacetate of Cyclosporin ene-yne-ene

[0157] To a 25 mL flask charged with a solution of the acetate of cyclosporin alkynyl alcohol from Example 86 (54 mg, 0.039 mmol) in benzene (2 mL) was added Burgess reagent (19 mg, 0.078 mmol). The resulting mixture was heated at 60°C. After 3.5 h, an additional 8 mg of Burgess reagent were added. The reaction was again heated at 60°C for 2 h. The mixture was cooled to room temperature and partitioned between ether (60 mL) and water (15 mL). The aqueous layer was

extracted once with ether (30 mL). The combined ethereal layers were washed with brine (20 mL), dried over anhydrous sodium sulfate, and then concentrated under reduced pressure. The crude material was purified by semi-preparative HPLC to yield the diacetate of cyclosporin ene-yne-ene (18 mg, 35%): ^1H NMR (300 MHz, CDCl_3) δ 8.54 (d, $J = 9.7$ Hz, 1H), 8.03 (d, $J = 6.8$ Hz, 1H), 7.62 (d, $J = 7.7$ Hz, 1H), 7.51 (d, $J = 9.1$ Hz, 1H), 6.04–5.86 (m, 1H), 5.74–5.65 (m, 1H), 5.88 (dd, $J = 17.5$, 2.2 Hz, 2H), 5.57–5.42 (m, 3H), 4.95 (t, $J = 7.5$ Hz, 1H), 5.19–4.93 (m, 3H), 4.85 (t, $J = 7.4$ Hz, 1H), 4.79 (t, $J = 12.0$ Hz, 1H), 4.57–3.31 (m, 3H), 3.37 (s, 3H), 3.27 (s, 3H), 3.25 (s, 3H), 3.20 (s, 3H), 3.13 (s, 3H), 2.68 (s, 3H), 2.67 (s, 3H), 2.47–2.31 (m, 1H), 2.24–2.10 (m, 3H), 2.10 (s, 3H), 2.02 (s, 3H), 1.98–1.82 (m, 4H), 1.80–1.60 (m, 4H), 1.48–0.72 (m, 58H); ESI MS m/z 1353 [$\text{C}_{70}\text{H}_{117}\text{N}_{11}\text{O}_{15} + \text{H}$] $^+$.

Example 89 - Preparation of Cyclosporin ene-yne-ene

[0158] A solution of the diacetate of cyclosporin ene-yne-ene from Example 88 (28 mg, 0.02 mmol) in methanol (2 mL) was treated with potassium carbonate (55 mg, 0.40 mmol). This was allowed to stir at room temperature overnight, after which it was quenched with a saturated solution of ammonium chloride (10 mL). The aqueous layer was then extracted with ethyl acetate (2 \times 40 mL). The combined organics were washed with brine (15 mL), dried over anhydrous sodium sulfate, and then concentrated under reduced pressure. The crude material was purified by semi-preparative HPLC to yield cyclosporin ene-yne-ene (8 mg, 32%) as a white solid: ^1H NMR (300 MHz, CDCl_3) δ 8.07 (d, $J = 9.8$ Hz, 1H), 7.70 (d, $J = 7.3$ Hz, 1H), 7.58 (d, $J = 8.0$ Hz, 1H), 7.27 (d, $J = 8.4$ Hz, 1H), 6.14–6.04 (m, 1H), 5.97–5.85 (m, 1H), 5.76–5.66 (m, 1H), 5.63 (d, $J = 2.0$ Hz, 1H), 5.58–5.41 (m, 4H), 5.30 (dd, $J = 11.4$, 3.1 Hz, 1H), 5.17–4.91 (m, 6H), 4.83 (t, $J = 7.1$ Hz, 1H), 4.65 (t, $J = 9.1$ Hz, 1H), 4.55 (t, $J = 7.2$ Hz, 1H), 4.05 (d, $J = 6.6$ Hz, 1H), 3.73 (t, $J = 6.6$ Hz, 1H), 3.50 (s, 3H), 3.30 (s, 3H), 3.26 (s, 3H), 3.15 (s, 3H), 3.10 (s, 3H), 2.71 (s, 3H), 2.69 (s, 3H), 2.63–2.52 (m, 1H), 2.44–2.34 (m, 1H), 2.23–1.88 (m, 5H), 1.83–1.60 (m, 7H), 1.50–0.69 (m, 54H); ESI MS m/z 1269 [$\text{C}_{66}\text{H}_{113}\text{N}_{11}\text{O}_{13} + \text{H}$] $^+$; HPLC 92.7% (AUC), $t_R = 19.67$ min.

Example 90 - Preparation of the Diacetate of Cyclosporin Alkynyl Bromide

[0159] To a flask charged with a solution of the acetate of cyclosporin alkyne from Example 78 (33 mg, 0.02 mmol) in acetone (2 mL) was added silver nitrate (3.4 mg, 0.02 mmol) and recrystallized *N*-bromosuccinimide (36 mg, 0.2 mmol). After 2 h, the reaction was cooled to 0°C and 5 mL of water were added. The aqueous layer was then extracted with ether (3 × 75 mL). The combined ethereal layers were washed with water (20 mL) and then brine (20 mL). The organics were dried over anhydrous sodium sulfate and then concentrated under reduced pressure.

10 The crude material was purified by semi-preparative HPLC to yield the diacetate of cyclosporin alkynyl bromide (23 mg, 80%): ¹H NMR (300 MHz, CDCl₃) δ 8.54 (d, *J* = 9.7 Hz, 1H), 8.02 (d, *J* = 6.7 Hz, 1H), 7.67 (d, *J* = 7.7 Hz, 1H), 7.57 (d, *J* = 9.1 Hz, 1H), 6.11–6.00 (m, 1H), 5.69 (dd, *J* = 11.1, 4.0 Hz, 1H), 5.58–5.42 (m, 1H), 5.40–5.13 (m, 5H), 5.07–4.91 (m, 3H), 4.88 (t, *J* = 7.1 Hz, 1H), 4.85 (t, *J* = 9.8 Hz, 1H), 4.57–4.34 (m, 3H), 3.37 (s, 3H), 3.26 (s, 3H), 3.24 (s, 3H), 3.19 (s, 3H), 2.68 (s, 6H), 2.48–2.32 (m, 1H), 2.27–2.12 (m, 3H), 2.11 (s, 3H), 2.02 (s, 3H), 1.98–1.61 (m, 7H), 1.49–0.66 (m, 59H); ESI MS *m/z* 1406 [C₆₈H₁₁₄BrN₁₁O₁₅ + H]⁺.

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Example 91 - Preparation of Cyclosporin Alkynyl Bromide

20 [0160] A solution of the diacetate of cyclosporin alkynyl bromide from Example 90 (22.5 mg, 0.02 mmol) in methanol (2 mL) was treated with potassium carbonate (55 mg, 0.40 mmol). The resulting mixture was stirred at room temperature for 4 h, after which it was quenched with a saturated solution of ammonium chloride (10 mL). The aqueous layer was then extracted with ethyl acetate (3 × 30 mL). The combined organics were washed with brine (20 mL), dried over anhydrous sodium sulfate, and then concentrated under reduced pressure. The crude material was purified by semi-preparative HPLC to yield cyclosporin alkynyl bromide (4.7 mg, 18%): ¹H NMR (300 MHz, CDCl₃) δ 8.09 (d, *J* = 10.0 Hz, 1H), 7.70 (d, *J* = 7.4 Hz, 1H), 7.54 (d, *J* = 8.3 Hz, 1H), 7.27 (d, *J* = 9.9 Hz, 1H), 6.24–6.21 (m, 1H), 5.70 (dd, *J* = 11.0, 3.9 Hz, 1H), 5.49 (d, *J* = 6.6 Hz, 1H), 5.44–5.34 (m, 1H), 5.30 (dd, *J* = 11.8, 3.8 Hz, 1H), 5.13–4.92 (m, 5H), 4.80 (t, *J* = 5.6 Hz, 1H), 4.65 (t, *J* = 8.6 Hz, 1H), 4.51 (t, *J* = 6.8 Hz, 1H), 4.04 (d, *J* = 6.7 Hz, 2H), 3.77 (t, *J* = 6.4 Hz, 1H), 3.49 (s, 3H), 3.30 (s, 3H), 3.26 (s, 3H), 3.15 (s, 3H), 3.11 (s, 3H), 2.70 (s, 3H), 2.69 (s, 3H), 2.57–

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2.36 (m, 2H), 2.21–1.95 (m, 6H), 1.85–1.56 (m, 7H), 1.50–0.72 (m, 54H); ESI MS m/z 1320 [$C_{64}H_{110}BrN_{11}O_{13} + H$] $^{+}$; HPLC >99% (AUC), t_R = 19.68 min.

Example 92 - Preparation of the Diacetate of Cyclosporin Alkyne

5 **[0161]** A flask charged with a solution of cyclopropyl(trimethylsilyl)acetylene (0.1 mL, 0.6 mmol) in triethylamine (3 mL) was cooled to 0°C. Tetrabutylammonium fluoride (0.72 mL, 1 M in THF, 0.72 mmol) was added and the mixture was allowed to stir for 10 min. The ice bath was removed and the flask warmed to room
10 temperature over 30 min. The acetate of *trans*-cyclosporin vinyl iodide from Example 77 (80 mg, 0.06 mmol) was added followed by copper(I) iodide (5.7 mg, 0.03 mmol) and dichlorobis(triphenylphosphine)palladium(II) (21 mg, 0.03 mmol). After 2 h, the reaction mixture was filtered twice through a microfilter then concentrated under reduced pressure. The crude material was purified by semi-
15 preparative HPLC to afford the diacetate of cyclosporin alkyne (47 mg, 57%): 1H NMR (300 MHz, $CDCl_3$) δ 8.54 (d, J = 9.5 Hz, 1H), 8.04 (d, J = 6.8 Hz, 1H), 7.66 (d, J = 7.7 Hz, 1H), 7.57 (d, J = 9.1 Hz, 1H), 5.89–5.77 (m, 1H), 5.69 (dd, J = 10.9, 3.6 Hz, 1H), 5.57–5.34 (m, 3H), 5.34–5.13 (m, 3H), 5.08–4.94 (m, 3H), 4.86 (t, J = 7.1 Hz, 1H), 4.79 (t, J = 9.3 Hz, 1H), 4.57–4.33 (m, 3H), 3.37 (s, 3H), 3.26 (s, 3H),
20 3.23 (s, 3H), 3.19 (s, 3H), 3.13 (s, 3H), 2.68 (s, 6H), 2.44–2.31 (m, 1H), 2.27–2.14 (m, 2H), 2.15 (s, 3H), 2.01 (s, 3H), 1.93–1.63 (m, 7H), 1.57–0.63 (m, 62H); ESI MS m/z 1367 [$C_{71}H_{119}N_{11}O_{15} + H$] $^{+}$.

Example 93 - Preparation of Cyclosporin Alkyne

25 **[0162]** A solution of the diacetate of cyclosporin alkyne from Example 92 (47 mg, 0.03 mmol) in methanol (2 mL) was treated with potassium carbonate (82 mg, 0.60 mmol). The resulting mixture was stirred at room temperature for 16 h, after which it was quenched with a saturated solution of ammonium chloride (10 mL).
30 The aqueous layer was then extracted with ethyl acetate (3 \times 75 mL). The combined organics were washed with brine (15 mL), dried over anhydrous sodium sulfate, and then concentrated under reduced pressure. The crude material was purified by semi-preparative HPLC to yield cyclosporin alkyne (25 mg, 66%) as a white solid: 1H NMR (300 MHz, $CDCl_3$) δ 8.04 (d, J = 10.1 Hz, 1H), 7.66 (d, J = 7.7 Hz, 1H), 7.44

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(d, $J = 8.3$ Hz, 1H), 7.16 (d, $J = 7.9$ Hz, 1H), 6.04–5.92 (m, 1H), 5.70 (dd, $J = 10.9$, 3.6 Hz, 1H), 5.51 (d, $J = 6.2$ Hz, 1H), 5.38–5.31 (m, 1H), 5.28 (dd, $J = 12.2$, 4.0 Hz, 1H), 5.13–4.89 (m, 5H), 4.83 (t, $J = 7.2$ Hz, 1H), 4.64 (t, $J = 9.7$ Hz, 1H), 4.54 (t, $J = 7.5$ Hz, 1H), 4.03 (d, $J = 6.7$ Hz, 2H), 3.70 (t, $J = 6.2$ Hz, 1H), 3.49 (s, 3H), 3.30 (s, 3H), 3.27 (s, 3H), 3.14 (s, 3H), 2.70 (s, 3H), 2.68 (s, 3H), 2.59–2.30 (m, 2H), 2.23–1.95 (m, 6H), 1.85–1.56 (m, 7H), 1.50–0.66 (m, 65H); ESI MS m/z 1283 $[\text{C}_{67}\text{H}_{115}\text{N}_{11}\text{O}_{13} + \text{H}]^+$; HPLC 97.7% (AUC), $t_R = 19.58$ min.

Example 94 - Preparation of the Acetate of Cyclosporin Alkynyl Thioether

[0163] To a flask charged with a solution of the acetate of *trans*-cyclosporin vinyl iodide from Example 77 (70 mg, 0.05 mmol) in triethylamine (3 mL) was added copper(I) iodide (5.7 mg, 0.03 mmol) followed by dichlorobis(triphenylphosphine)palladium(II) (21 mg, 0.03 mmol) and propargyl ethyl sulphide (0.1 mL, 0.5 mmol). After 5.5 h, an additional 21 mg of dichlorobis(triphenylphosphine)palladium(II) were added and the reaction was left to stir overnight. The reaction mixture was filtered through a microfilter then concentrated under reduced pressure. The crude material was purified by semi-preparative HPLC to afford the acetate of cyclosporin alkynyl thioether (45 mg, 64%): ^1H NMR (300 MHz, CDCl_3) δ 8.54 (d, $J = 9.7$ Hz, 1H), 8.04 (d, $J = 6.8$ Hz, 1H), 7.67 (d, $J = 7.7$ Hz, 1H), 7.62 (d, $J = 9.1$ Hz, 1H), 5.96–5.84 (m, 1H), 5.70 (dd, $J = 11.1$, 3.9 Hz, 1H), 5.57–5.42 (m, 2H), 5.39–5.13 (m, 5H), 5.08–4.97 (m, 3H), 4.86 (t, $J = 7.2$ Hz, 1H), 4.78 (t, $J = 9.6$ Hz, 1H), 4.59–4.33 (m, 3H), 3.37 (s, 3H), 3.26 (s, 3H), 3.23 (s, 3H), 3.19 (s, 3H), 3.13 (s, 3H), 2.69 (s, 3H), 2.68 (s, 3H), 2.45–2.34 (m, 1H), 2.24–2.12 (m, 2H), 2.14 (s, 3H), 2.02 (s, 3H), 1.97–1.83 (m, 5H), 1.76–1.13 (m, 4H), 1.48–0.75 (m, 61H); ESI MS m/z 1401 $[\text{C}_{71}\text{H}_{121}\text{N}_{11}\text{O}_{15}\text{S} + \text{H}]^+$.

Example 95 - Preparation of Cyclosporin Alkynyl Thioether

[0164] A solution of the acetate of cyclosporin alkynyl thioether from Example 94 (45 mg, 0.03 mmol) in methanol (2 mL) was treated with potassium carbonate (83 mg, 0.60 mmol). The resulting mixture was stirred at room temperature for 16 h, after which it was quenched with a saturated solution of ammonium chloride (10 mL). The aqueous layer was then extracted with ethyl acetate (3×75 mL). The

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combined organics were washed with brine (50 mL), dried over anhydrous sodium sulfate, and then concentrated under reduced pressure. The crude material was purified by semi-preparative HPLC to yield cyclosporin alkynyl thioether (23 mg, 58%) as a white solid: ^1H NMR (300 MHz, CDCl_3) δ 8.04 (d, J = 10.0 Hz, 1H), 7.66 (d, J = 7.8 Hz, 1H), 7.45 (d, J = 8.2 Hz, 1H), 7.15 (d, J = 7.8 Hz, 1H), 6.13–5.99 (m, 1H), 5.70 (dd, J = 11.1, 4.2 Hz, 1H), 5.53 (d, J = 5.9 Hz, 1H), 5.45–5.36 (m, 1H), 5.28 (dd, J = 11.7, 3.8 Hz, 1H), 5.14–4.87 (m, 3H), 4.83 (t, J = 7.3 Hz, 1H), 4.64 (t, J = 9.5 Hz, 1H), 4.55 (t, J = 7.5 Hz, 1H), 4.04 (d, J = 6.8 Hz, 2H), 3.70 (t, J = 6.3 Hz, 1H), 3.50 (s, 3H), 3.40 (s, 3H), 3.30 (s, 3H), 3.27 (s, 3H), 3.15 (s, 3H), 2.73–2.64 (m, 9H), 2.61–2.52 (m, 1H), 2.45–2.34 (m, 1H), 2.24–1.89 (m, 12H), 1.83–1.57 (m, 8H), 1.48–0.75 (m, 53H); ESI MS m/z 1317 [$\text{C}_{67}\text{H}_{117}\text{N}_{11}\text{O}_{13}\text{S} + \text{H}$] $^+$; HPLC 94.0% (AUC), t_R = 19.80 min.

Example 96 - Preparation of the Acetate of Cyclosporin Alkynyl Ether

[0165] To a flask charged with a solution of the acetate of *trans*-cyclosporin vinyl iodide from Example 77 (70 mg, 0.05 mmol) in triethylamine (3 mL) was added copper(I) iodide (5.7 mg, 0.03 mmol) followed by dichlorobis(triphenylphosphine)palladium(II) (21 mg, 0.03 mmol) and methyl propargyl ether (0.05 mL, 0.5 mmol). After 6 h, the reaction mixture was filtered through a microfilter then concentrated under reduced pressure. The crude material was purified by semi-preparative HPLC to afford the acetate of cyclosporin alkynyl ether (52 mg, 75%): ^1H NMR (300 MHz, CDCl_3) δ 8.52 (d, J = 9.6 Hz, 1H), 8.01 (d, J = 6.8 Hz, 1H), 7.59 (d, J = 7.7 Hz, 1H), 7.46 (d, J = 9.1 Hz, 1H), 6.06–5.93 (m, 1H), 5.68 (dd, J = 10.9, 3.8 Hz, 1H), 5.57–5.42 (m, 2H), 5.37–5.31 (m, 2H), 5.08–4.97 (m, 3H), 5.29 (dd, J = 8.9, 2.6 Hz, 1H), 5.15 (t, J = 5.6 Hz, 1H), 5.19–4.92 (m, 3H), 4.85 (t, J = 7.2 Hz, 1H), 4.78 (t, J = 9.6 Hz, 1H), 3.39 (s, 3H), 3.37 (s, 3H), 3.27 (s, 3H), 3.24 (s, 3H), 3.20 (s, 3H), 3.18 (s, 3H), 2.68 (s, 3H), 2.67 (s, 3H), 2.47–2.33 (m, 1H), 2.33–2.13 (m, 3H), 2.10 (s, 3H), 2.10 (s, 3H), 1.95–1.83 (m, 4H), 1.43–0.75 (m, 61H); ESI MS m/z 1371 [$\text{C}_{70}\text{H}_{119}\text{N}_{11}\text{O}_{16} + \text{H}$] $^+$.

Example 97 - Preparation of Cyclosporin Alkynyl Ether

[0166] A solution of the acetate of cyclosporin alkynyl ether from Example 96 (52 mg, 0.04 mmol) in methanol (2 mL) was treated with potassium carbonate (111 mg, 0.80 mmol). The resulting mixture was stirred at room temperature for 16 h, after which it was quenched with a saturated solution of ammonium chloride (10 mL). The aqueous layer was then extracted with ethyl acetate (3 × 75 mL). The combined organics were washed with brine (15 mL), dried over anhydrous sodium sulfate, and then concentrated under reduced pressure. The crude material was purified by semi-preparative HPLC to yield cyclosporin alkynyl ether (37 mg, 73%) as a white solid: ¹H NMR (300 MHz, CDCl₃) δ 8.07 (d, *J* = 9.8 Hz, 1H), 7.67 (d, *J* = 7.6 Hz, 1H), 7.46 (d, *J* = 8.2 Hz, 1H), 7.17 (d, *J* = 8.0 Hz, 1H), 6.19–6.07 (m, 1H), 5.70 (dd, *J* = 10.9, 3.9 Hz, 1H), 5.52 (d, *J* = 6.1 Hz, 1H), 5.49–5.41 (m, 1H), 5.27 (dd, *J* = 11.7, 3.8 Hz, 1H), 5.11–4.91 (m, 6H), 4.83 (t, *J* = 7.4 Hz, 1H), 4.64 (t, *J* = 8.5 Hz, 1H), 4.54 (t, *J* = 7.4 Hz, 1H), 4.20 (d, *J* = 1.8 Hz, 2H), 4.04 (d, *J* = 6.8 Hz, 2H), 3.72 (t, *J* = 6.6 Hz, 1H), 3.50 (s, 3H), 3.39 (s, 3H), 3.30 (s, 3H), 3.27 (s, 3H), 3.15 (s, 3H), 3.11 (s, 3H), 2.70 (s, 3H), 2.68 (s, 3H), 2.53–2.41 (m, 5H), 2.41–2.21 (m, 6H), 2.20–1.92 (m, 6H), 1.85–1.56 (m, 7H), 1.51–0.66 (m, 44H); ESI MS *m/z* 1287 [C₆₆H₁₁₅N₁₁O₁₄ + H]⁺; HPLC >99% (AUC), *t_R* = 18.42 min.

Example 98 - Concanavalin A- Stimulated Splenocyte Assay

[0167] Male BALB/c mice, at 5 to 7 weeks of age, were sacrificed by CO₂ inhalation. Spleens were removed and dissociated by pushing through a nylon cell strainer. The splenocytes were washed in RPMI 1640/5% fetal calf serum (FCS) and pelleted at 400×g. Red blood cells were then lysed by resuspending the cell pellet in ACK lysis buffer (150 mM NH₄Cl, 1 mM KHCO₃, 0.1 mM EDTA, 3 mL per spleen) for 10 min at room temperature. After pelleting at 400×g, the cells were washed by resuspending in RPMI 1640/5% FCS and repelleting. The cell pellet was resuspended in RPMI 1640/5% FCS and again passed through a cell strainer to remove cell aggregates. The cells were then counted and adjusted to 2 × 10⁶ cells/ml in RPMI 1640/10% FCS/50 μM 2-mercaptoethanol. Cell viability was assessed by Trypan blue staining. Cyclosporin A or the test compound and two micrograms of concanavalin A were added to the wells of a 96 well plate, prior to the addition of 2 ×

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10⁵ splenocytes. The cells were cultured in a 37°C CO₂ incubator for 2 days and then pulsed with 1 µCi of [³H]thymidine for 6 hours. Cells were harvested onto filtermats with a TomTec 96 well plate harvester and lysed with H₂O. The filtermat and scintillation fluid were sealed in a plastic sleeve. [³H]thymidine incorporation was measured with a Wallac Trilux plate counter. Initial screens were done at a fixed value of 100 ng/ml test compound. IC₅₀s were calculated from 7 point concentration-response curves using GraphPad software.

Example 99 - Murine Ex Vivo Pharmacodynamic Assay

10 [0168] *In vivo* immunosuppressive activity can be determined for cyclosporin A and the disclosed cyclosporin analogs, as described below. The concanavalin A-stimulated splenocyte activity can be assessed *in vivo* using a method previously described by Peterson et al. (Peterson et al., "A Tacrolimus-Related

15 Immunosuppressant with Biochemical Properties Distinct from Those of Tacrolimus," *Transplantation*, 65:10-18 (1998), which is hereby incorporated by reference in its entirety) or a slightly modified version thereof.

[0169] Optimal doses of cyclosporin A or an immunosuppressive compound of the present invention (four different doses of test drug plus a control set of animals

20 with no drug) was administered orally or intravenously to male BALB/c or female C57BL mice. Three mice were tested at each dose. Concanavalin A was injected into the tail vein of the mouse at 4 hours after the administration of cyclosporin A or the immunosuppressive compound. One hour after the concanavalin A injection, the mice were euthanized, the spleens were removed under sterile conditions, and the

25 extent of splenocyte proliferation was measured in a similar manner as described in Example 98. The percent inhibition relative to control was plotted graphically versus the dose of the immunosuppressive compound and an ED₅₀ value was determined. Each dose-response assay for the compound of the present invention was accompanied by a cyclosporin control at a single dose equal to the ED₅₀.

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Example 100 - Assay for Inhibition of Peptidyl Prolyl Isomerase Activity of Cyclophilin A

[0170] The assay for inhibition of peptidyl prolyl isomerase activity of
5 cyclophilin A is a modification of the procedure described by Kofron et al.,
“Determination of Kinetic Constants for Peptidyl Prolyl *cis-trans* Isomerases by an
Improved Spectrophotometric Assay,” *Biochemistry* 30:6127-6134 (1991), which is
hereby incorporated by reference in its entirety. Recombinant human cyclophilin A in
50 mM HEPES, 100 mM NaCl pH 8.0 is precooled to 4°C. Test compounds and the
10 cyclosporin positive control are dissolved in dimethyl sulfoxide (DMSO) and
introduced over a range of concentrations. Chymotrypsin is then added to a final
concentration of 6 mg/ml. The peptide substrate, Suc-Ala-Ala-Pro-Phe-pNA (SEQ ID
NO: 1), is dissolved in 470 mM LiCl in trifluoroethanol and then added to 25 µg/ml to
initiate the reaction. After rapid mixing, the absorbance at 390 nm is monitored over
15 a 90 second time course.

Example 101 - Cellular Assay for Determination of HIV Inhibition

[0171] The *in vitro* anti-HIV activity of compounds of the present invention is
20 measured in established cell line cultures as described by Mayaux et al., “Triterpene
Derivatives That Block Entry of Human Immunodeficiency Virus Type 1 Into Cells,”
Proc. Natl. Acad. Sci. USA 91:3564-3568 (1994), which is hereby incorporated by
reference in its entirety. The CEM4 cell line was infected with HIV-1_{Lai} strain. The
inhibition of HIV replication in the culture is estimated by the measure of the reverse
25 transcriptase (RT) produced in the supernatant. Anti-viral activity is expressed as the
IC₅₀ RT, the concentration required to reduce replication of HIV by 50%, and is
determined by linear regression.

Example 102 – Intracellular Replication of the HCV Genome *in vitro*

30 [0172] The effect of the cyclosporin compounds of the present invention on
the intracellular replication of the HCV genome *in vitro*, using an HCV replicon
system in a cultured human hepatoma Huh7 cell line is determined by the method of
Lohmann et al., “Replication of Subgenomic Hepatitis C Virus RNAs in a Hepatoma

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Cell Line,” *Science* 285:110-113 (1999), which is hereby incorporated by reference in its entirety.

Example 103 –In vitro HCV Infection Experiment

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[0173] The *in vitro* HCV infection experiment is performed as described by Kato et al., “Replication of Hepatitis C Virus in Cultured Non-Neoplastic Human Hepatocytes,” *Jpn. J. Cancer Res.* 87:787-792 (1996), which is hereby incorporated by reference in its entirety, and Ikada et al., “Human Hepatocyte Clonal Cell Lines That Support Persistent Replication of Hepatitis C Virus,” *Virus Res.* 56:157-167 (1998), which is hereby incorporated by reference in its entirety.

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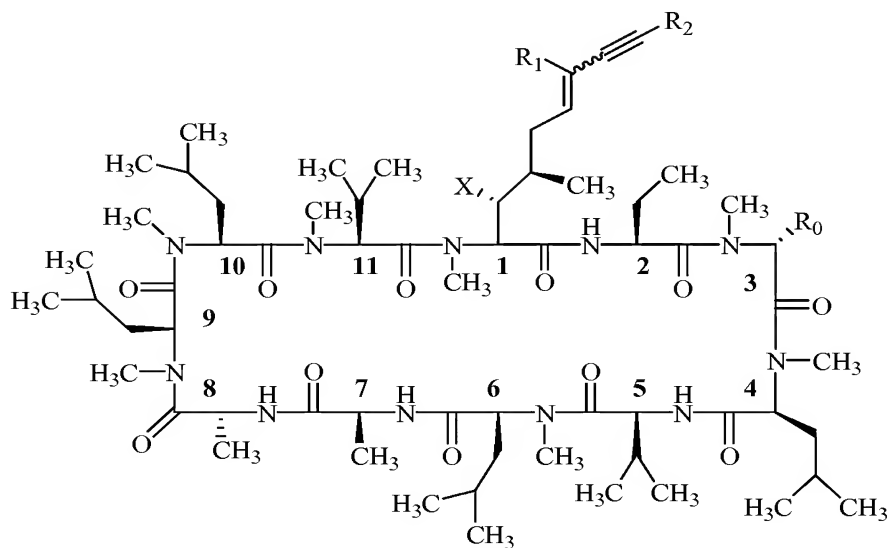
[0174] Although the invention has been described in detail for the purpose of illustration, it is understood that such detail is solely for that purpose, and variations can be made therein by those skilled in the art without departing from the spirit and scope of the invention which is defined by the following claims.

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WHAT IS CLAIMED IS:

1. A method of preventing or treating a mammal with a viral-induced disorder comprising:

administering to the mammal a therapeutically effective amount of a compound having the following formula:

**Formula I**

wherein:

X is OH or OAc;

R₀ is H, CH₂OH, or CH₂OR₃;

R₁ is hydrogen, deuterium, or methyl;

R₂ is selected from the group consisting of:

hydrogen;

halogen;

C₁-C₆ saturated or unsaturated, straight or branched carbon chain;

C₁-C₆ saturated or unsaturated, straight or branched carbon chain containing a substitution or substitutions selected from the group consisting of deuterium, halogen, nitrogen, sulfur, and silicon;

C₁-C₆ saturated or unsaturated, straight or branched carbon chain containing a function group or function groups selected from the group consisting of alcohol, ether, aldehyde, ketone, carboxylic acid, ester, and amide;

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C₁-C₆ saturated or unsaturated, straight or branched carbon chain containing a function group of oxime or hydrazone;

C₁-C₆ saturated or unsaturated, straight or branched carbon chain containing an aryl or a heteroaryl group;

C₃-C₆ substituted and unsubstituted cycloalkyl;
substituted and unsubstituted aryl; and
substituted and unsubstituted heteroaryl; and

R₃ is selected from the group consisting of:

alkanoyl,
alkenoyl,
alkynoyl,
aryloyl,
arylalkanoyl,
alkylaminocarbonyl,
arylaminocarbonyl,
arylalkylaminocarbonyl,
alkyloxycarbonyl,
aryloxycarbonyl, and
arylalkyloxycarbonyl,

wherein the compound is a cis geometric isomer, a trans geometric isomer, or a mixture of the cis and the trans geometric isomers or a pharmaceutically acceptable salt thereof,

under conditions effective to prevent or treat the viral-induced disorder.

2. The method according to claim 1, wherein X is OH or OAc, R₀ is H, CH₂OH, or CH₂OAc, and R₁ is H or D.

3. The method according to claim 2, wherein R₂ is H.

4. The method according to claim 2, wherein R₂ is selected from the group consisting of CH₃, CD₃, CH₂CH₃, and CH₂CH₂CH₃.

5. The method according to claim 2, wherein R₂ is selected from the group consisting of CH=CH₂, CH=CHCH₃, C≡CH, and -C≡C-CH₃.

6. The method according to claim 2, wherein R₂ is selected from the group consisting of F, Cl, Br, and I.

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7. The method according to claim 2, wherein R₂ is cyclopropyl.
8. The method according to claim 2, wherein R₂ is selected from the group consisting of CH₂OH, CH(OH)CH₃, CH₂OCH₃, CH₂OCH₂CH₃, CHO, and C(=O)CH₃.
9. The method according to claim 2, wherein R₂ is selected from the group consisting of CH=N-OCH₃, CH=N-OCH₂CH₃, CH=N-NHCH₃, and CH=N-N(CH₃)₂.
10. The method according to claim 1, wherein the viral-induced disorder is a human immunodeficiency virus-induced disorder.
11. The method according to claim 10, wherein said compound is administered in combination with antiretroviral agents selected from the group consisting of nucleoside reverse transcriptase inhibitors, nonnucleoside reverse transcriptase inhibitors, human immunodeficiency virus protease inhibitors, fusion inhibitors, and combinations thereof.
12. The method according to claim 11, wherein the nucleoside reverse transcriptase inhibitor is selected from the group consisting of Zidovudine, Didanosine, Stavudine, and Lamivudine.
13. The method according to claim 11, wherein the nonnucleoside reverse transcriptase inhibitor is selected from the group consisting of Nevirapine, Efavirenz, and Delavirdine.
14. The method according to claim 11, wherein the human immunodeficiency virus protease inhibitor is selected from the group consisting of Saquinovir, Indinavir, and Ritonavir.
15. The method according to claim 11, wherein the fusion inhibitor is Enfuvirtide.

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16. The method according to claim 1, wherein the viral-induced disorder is a hepatitis C virus-induced disorder.

17. The method according to claim 16, wherein said compound is administered in combination with an interferon.

18. The method according to claim 17, wherein the interferon is interferon α 2a or interferon α 2b.

19. The method according to claim 17, wherein the interferon is a pegylated interferon.

20. The method according to claim 19, wherein the pegylated interferon is pegylated interferon α 2a or pegylated interferon α 2b.

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Figure 1

